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FILE LAST UPDATED: 16 Nov 2008 (20081116/ED)

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L1 ANSWER 1 OF 1 CAPLUS COPYRIGHT 2008 ACS on STN  
 AN 2005:673860 CAPLUS  
 DN 143:176223  
 ED Entered STN: 31 Jul 2005  
 TI Composite mixed oxides as active battery electrodes, especially for rechargeable lithium batteries  
 IN Gauthier, Gilles; Le Cras, Frederic; Lignier, Helene; Gabelle, Jean Louis  
 PA Commissariat a l'Energie Atomique, Fr.  
 SO Fr. Demande, 45 pp.  
 CODEN: FRXXBL  
 DT Patent  
 LA French  
 IC ICM H01M004-60  
 ICS H01M004-26  
 CC 52-2 (Electrochemical, Radiational, and Thermal Energy Technology)  
 FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	FR 2865576	A1	20050729	FR 2004-50156	20040128
	FR 2865576	B1	20060428		
	WO 2005076390	A2	20050818	WO 2005-FR50045	20050126
	WO 2005076390	A3	20051006		
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	EP 1709702	A2	20061011	EP 2005-717687	20050126
	EP 1709702	B1	20071205		
	R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, FI, RO, CY, TR, BG, CZ, EE, HU, PL, SK, IS				
	JP 2007520038	T	20070719	JP 2006-550260	20050126
	AT 380399	T	20071215	AT 2005-717687	20050126
	ES 2299019	T3	20080516	ES 2005-717687	20050126
	US 20070152185	A1	20070705	US 2007-586601	20070104 <--
PRAI	FR 2004-50156	A	20040128		
	WO 2005-FR50045	W	20050126		

#### CLASS

	PATENT NO.	CLASS	PATENT FAMILY CLASSIFICATION CODES
	FR 2865576	ICM	H01M004-60
		ICS	H01M004-26
		IPCI	H01M0004-36 [I,C]; H01M0004-26 [I,C]; H01M0004-60 [I,A]; H01M0004-26 [I,A]
		IPCR	H01M0004-36 [I,C]; H01M0004-60 [I,A]; C01G0001-02 [I,C*]; C01G0001-02 [I,A]; C01G0001-06 [I,C*]; C01G0001-06 [I,A]; H01M0004-26 [I,C]; H01M0004-26 [I,A]; H01M0004-48 [I,C*]; H01M0004-48 [I,A]; H01M0004-50 [N,C*]; H01M0004-50 [N,A]; H01M0004-52 [N,C*]; H01M0004-52 [N,A]; H01M0004-58 [I,C*]; H01M0004-58 [I,A]; H01M0004-62 [I,C*]; H01M0004-62 [I,A]
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		H01M0004-58 [I,C*]; H01M0004-58 [I,A]; H01M0004-62 [I,C*]; H01M0004-62 [I,A]
EP 1709702	ECLA	C01G001/02; C01G001/06; H01M004/48B2; H01M004/62C2; M01P; T01M; T01M; T01M
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JP 2007520038	IPCI	H01M0004-04 [I,A]; H01M0004-58 [I,A]; H01M0004-62 [I,A]; H01M0010-40 [N,A]; H01M0010-36 [N,C*]
	IPCR	H01M0004-04 [I,C]; H01M0004-04 [I,A]; C01G0001-02 [I,C*]; C01G0001-02 [I,A]; C01G0001-06 [I,C*]; C01G0001-06 [I,A]; H01M0004-48 [I,C*]; H01M0004-48 [I,A]; H01M0004-50 [N,C*]; H01M0004-50 [N,A]; H01M0004-52 [N,C*]; H01M0004-52 [N,A]; H01M0004-58 [I,C]; H01M0004-58 [I,A]; H01M0004-62 [I,C]; H01M0004-62 [I,A]; H01M0010-36 [N,C]; H01M0010-40 [N,A]
	ECLA	C01G001/02; C01G001/06; H01M004/48B2; H01M004/62C2; M01P; T01M; T01M
	FTERM	5H029/AJ03; 5H029/AJ12; 5H029/AJ14; 5H029/AK03; 5H029/AL06; 5H029/AL07; 5H029/AL08; 5H029/CJ02; 5H029/CJ08; 5H029/CJ28; 5H029/DJ08; 5H029/EJ04; 5H029/EJ11; 5H029/EJ12; 5H029/HJ01; 5H029/HJ02; 5H029/HJ05; 5H029/HJ07; 5H029/HJ14; 5H050/AA08; 5H050/AA15; 5H050/AA19; 5H050/BA16; 5H050/BA17; 5H050/CA07; 5H050/CA08; 5H050/CA09; 5H050/CB07; 5H050/CB08; 5H050/CB09; 5H050/DA09; 5H050/EA08; 5H050/EA22; 5H050/EA23; 5H050/GA02; 5H050/GA10; 5H050/GA27; 5H050/HA01; 5H050/HA02; 5H050/HA05; 5H050/HA07; 5H050/HA14; 5H050/HA20
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ES 2299019	IPCI	H01M0004-04 [I,C]; H01M0004-04 [I,A]; H01M0004-58 [I,C]; H01M0004-58 [I,A]
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	ECLA	C01G001/02; C01G001/06; H01M004/48B2; H01M004/62C2; M01P; T01M; T01M; T01M
US 20070152185	IPCI	H01B0001-06 [I,A]; H01B0001-18 [I,A]; H01B0001-14 [I,C*]
	NCL	252/182.100; 252/506.000; 252/507.000

AB Electrode-active materials, especially for alkali ion insertion (i.e., Na<sup>+</sup> and Li<sup>+</sup>) for lithium batteries, contain, as an active component, a composition of general formula AaDdMmZsOoNnFf,, in which: (1) A is an alkali metal, (2) D is an alkaline earth metal or a Group IIIA element, with the exception of B, (3) M is a transition metal, (4) Z is a non-metal selected from S, Se, P, As, Si, Ge, Sn, and B, (5) O is oxygen, N is nitrogen, and F is fluorine, and (6) a, d, m, z, o, n, and f are ≥0. The compns., which also contain an electron conductor, such as carbon, are prepared by thermal decomposition of homogeneously mixed precursors, which are organic or organometallic derivs. (preferably at 200–600°). Preferred components include: (1) A = Li, Na, and K, (2) D is Mg, Al, and Ga, (3) M = Fe, Ni, Co, Mn, V, Mo, Nb, W, and Ti; preferred components are LiFePO<sub>4</sub>, LiFeBO<sub>3</sub>, or NaFeBO<sub>3</sub>.

ST electrode mixed oxide lithium rechargeable battery; iron lithium borate secondary battery electrode

IT Transition metal oxides  
RL: DEV (Device component use); TEM (Technical or engineered material use); USES (Uses)  
(alkaline earth oxides, electrode active materials; composite mixed oxides as active battery electrodes, especially for rechargeable lithium batteries)

IT Transition metal oxides  
RL: DEV (Device component use); TEM (Technical or engineered material use); USES (Uses)  
(alkali metal oxides, electrode active materials; composite mixed oxides as active battery electrodes, especially for rechargeable lithium batteries)

IT Battery electrodes  
(composite mixed oxides as active battery electrodes, especially for rechargeable lithium batteries)

IT Carboxylic acids, processes  
RL: CPS (Chemical process); PEP (Physical, engineering or chemical process); PROC (Process)  
(dicarboxylic, metal salts and complexes, electrode precursors; thermal decomposition of; composite mixed oxides as active battery electrodes, especially for rechargeable lithium batteries)

IT Carboxylic acids, processes  
RL: CPS (Chemical process); PEP (Physical, engineering or chemical process); PROC (Process)  
(hydroxy, metal salts and complexes, electrode precursors; thermal decomposition of; composite mixed oxides as active battery electrodes, especially for rechargeable lithium batteries)

IT Amino acids, processes  
Polyoxyalkylenes, processes  
RL: CPS (Chemical process); PEP (Physical, engineering or chemical process); PROC (Process)  
(metal salts and complexes, electrode precursors; thermal decomposition of; composite mixed oxides as active battery electrodes, especially for rechargeable lithium batteries)

IT Carboxylic acids, processes  
RL: CPS (Chemical process); PEP (Physical, engineering or chemical process); PROC (Process)  
(oxo, metal salts and complexes, electrode precursors; thermal decomposition of; composite mixed oxides as active battery electrodes, especially for rechargeable lithium batteries)

IT Alkali metal oxides  
Alkaline earth oxides  
Group IIIA element oxides  
RL: DEV (Device component use); TEM (Technical or engineered material use); USES (Uses)  
(transition metal oxides, electrode active materials; composite mixed

oxides as active battery electrodes, especially for rechargeable lithium batteries)

IT 7440-44-0, Carbon, uses  
RL: TEM (Technical or engineered material use); USES (Uses)  
(elec. conductor; composite mixed oxides as active battery electrodes,  
especially for rechargeable lithium batteries)

IT 15365-14-7, Iron lithium phosphate (FeLiPO<sub>4</sub>) 332079-85-3, Iron lithium  
borate (FeLiBO<sub>3</sub>) 861001-97-0  
RL: DEV (Device component use); TEM (Technical or engineered material  
use); USES (Uses)  
(electrode active materials; composite mixed oxides as active battery  
electrodes, especially for rechargeable lithium batteries)

IT 50-21-5D, Lactic acid, metal salts and complexes 56-40-6D, Aminoacetic  
acid, metal salts and complexes 56-41-7D, Alanine, metal salts and  
complexes 56-84-8D, L-Aspartic acid, metal salts and complexes  
56-86-0D, L-Glutamic acid, metal salts and complexes 56-87-1D, L-Lysine,  
metal salts and complexes 61-90-5D, L-Leucine, metal salts and complexes  
70-26-8D, Ornithine, metal salts and complexes 74-79-3D, L-Arginine,  
metal salts and complexes 77-92-9D, Citric acid, metal salts and  
complexes 79-14-1D, Glycolic acid, metal salts and complexes 87-69-4D,  
Tartaric acid, metal salts and complexes 90-64-2D, Mandelic acid, metal  
salts and complexes 107-21-1D, Ethylene glycol, metal salts and  
complexes 110-15-6D, Succinic acid, metal salts and complexes  
110-16-7D, Maleic acid, metal salts and complexes 110-17-8D, Fumaric  
acid, metal salts and complexes 110-94-1D, Glutaric acid, metal salts  
and complexes 111-46-6D, Diethylene glycol, metal salts and complexes  
123-76-2D, Levulinic acid, metal salts and complexes 124-04-9D, Adipic  
acid, metal salts and complexes 127-17-3D, Pyruvic acid, metal salts and  
complexes 141-82-2D, Malonic acid, metal salts and complexes  
144-62-7D, Oxalic acid, metal salts and complexes 298-12-4D, Glyoxylic  
acid, metal salts and complexes 498-23-7D, Citraconic acid, metal salts  
and complexes 499-12-7D, Aconitic acid, metal salts and complexes  
6915-15-7D, Malic acid, metal salts and complexes 28854-76-4D, metal  
salts and complexes 35054-79-6D, Hydroxybutyric acid, metal salts and  
complexes 111937-70-3D, Hydroxyacrylic acid, metal salts and complexes  
151677-68-8  
RL: CPS (Chemical process); PEP (Physical, engineering or chemical  
process); PROC (Process)  
(electrode precursors; thermal decomposition of; composite mixed oxides as  
active battery electrodes, especially for rechargeable lithium batteries)

RE.CNT 7 THERE ARE 7 CITED REFERENCES AVAILABLE FOR THIS RECORD

RE

- (1) Commissariat Energie Atomique; CH 513769 A 1971 CAPLUS
- (2) Darmes, D; WO 0208355 A 2002 CAPLUS
- (3) Du, K; JOURNAL OF ALLOYS AND COMPOUNDS 2003, V352(1-2), P250 CAPLUS
- (4) Hydro Quebec; CA 2270771 A 2000 CAPLUS
- (5) L'Energie Atomique Et Institut Francais Du Petrole Des Carburants Et; BE 735476 A 1969 CAPLUS
- (6) Pechini, M; US 3330697 A 1967
- (7) Univ Texas; WO 9740541 A 1997 CAPLUS

=> s 15365-14-7 and 7440-44-0

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L5 1288 L4

L6 462 L5 AND L3

=> s 16 and electrode#  
730889 ELECTRODE#  
L7 155 L6 AND ELECTRODE#  
  
=> s 16 and (electrode# or anode# or cathode#)  
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184108 ANODE#  
227805 CATHODE#  
L8 432 L6 AND (ELECTRODE# OR ANODE# OR CATHODE#)

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70-26-8 or 74-79-3 or 77-92-9 or 79-14-1 or 87-69-4 or 90-64-2 or 107-21-1 or  
110-15-6 or 110-16-7 or 110-17-8

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L10 15717 L9

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Invalid automated search and crossover syntax. The information preceding the automated search and crossover field resulted in no query being identified. This is typically caused by the character preceding the "/" for the automated search and crossover field code representing a termination point, such as a left parenthesis "(".

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L14 76293 L13

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L18 47918 L17

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L20 65792 L19

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L22 62717 L21

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107-21-1 or 110-15-6 or 110-16-7 or 110-17-8

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L25 15717 L24

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L27 16848 L26

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L39        73205 L38

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L45        40764 L44

L46        258073 L45 OR L43 OR L41 OR L39 OR L37 OR L35 OR L33 OR L31 OR L29 OR  
L27 OR L25

=> s 110-94-1 or 111-46-6 or 123-76-2 or 124-04-9 or 127-17-3 or 141-82-2 or  
144-62-7 or 298-12-4 or 498-23-7 or 499-12-7

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L48            1047 L47

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L50            1043 L49

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L52            5548 L51

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L54            36542 L53

REGISTRY INITIATED  
Substance data SEARCH and crossover from CAS REGISTRY in progress...  
Use DISPLAY HITSTR (or FHITSTR) to directly view retrieved structures.

L56 14876 L55

REGISTRY INITIATED  
Substance data SEARCH and crossover from CAS REGISTRY in progress...  
Use DISPLAY HITSTR (or FHITSTR) to directly view retrieved structures.

L58 25465 L57

REGISTRY INITIATED  
Substance data SEARCH and crossover from CAS REGISTRY in progress...  
Use DISPLAY HITSTR (or FHITSTR) to directly view retrieved structures.

L60 16236 L59

REGISTRY INITIATED  
Substance data SEARCH and crossover from CAS REGISTRY in progress...  
Use DISPLAY HITSTR (or FHITSTR) to directly view retrieved structures.

L62 2735 L61

REGISTRY INITIATED  
Substance data SEARCH and crossover from CAS REGISTRY in progress...  
Use DISPLAY HITSTR (or FHITSTR) to directly view retrieved structures.

L64 15270 L63

REGISTRY INITIATED  
Substance data SEARCH and crossover from CAS REGISTRY in progress...  
Use DISPLAY HITSTR (or FHITSTR) to directly view retrieved structures.

L66 6189 L65

L67 108496 L66 OR L64 OR L62 OR L60 OR L58 OR L56 OR L54 OR L52 OR L50 OR  
L48

=> s 6915-15-7 or 28854-76-4 or 35054-79-6 or 111937-70-3 or 151677-68-8  
REG1stRY INITIATED  
Substance data SEARCH and crossover from CAS REGISTRY in progress...  
Use DISPLAY HITSTR (or FHITSTR) to directly view retrieved structures.

L69 7 L68

REG1stRY INITIATED  
Substance data SEARCH and crossover from CAS REGISTRY in progress...  
Use DISPLAY HITSTR (or FHITSTR) to directly view retrieved structures.

L71 32 L70

REG1stRY INITIATED  
Substance data SEARCH and crossover from CAS REGISTRY in progress...  
Use DISPLAY HITSTR (or FHITSTR) to directly view retrieved structures.

L73 286 L72

REG1stRY INITIATED  
Substance data SEARCH and crossover from CAS REGISTRY in progress...  
Use DISPLAY HITSTR (or FHITSTR) to directly view retrieved structures.

L75            74 L74

REGISTRY INITIATED  
Substance data SEARCH and crossover from CAS REGISTRY in progress...  
Use DISPLAY HITSTR (or FHITSTR) to directly view retrieved structures.

L77            24029 L76

L78            24346 L77 OR L75 OR L73 OR L71 OR L69

=> d his

(FILE 'HOME' ENTERED AT 14:46:54 ON 17 NOV 2008)

FILE 'CAPLUS' ENTERED AT 14:47:20 ON 17 NOV 2008  
E US20070152185/PN

L1            1 S E3  
S 15365-14-7/REG# AND 7440-44-0/REG#

FILE 'REGISTRY' ENTERED AT 14:48:47 ON 17 NOV 2008  
L2            1 S 7440-44-0/RN

FILE 'CAPLUS' ENTERED AT 14:48:47 ON 17 NOV 2008  
L3            387474 S L2

FILE 'REGISTRY' ENTERED AT 14:48:47 ON 17 NOV 2008  
L4            1 S 15365-14-7/RN

FILE 'CAPLUS' ENTERED AT 14:48:48 ON 17 NOV 2008  
L5            1288 S L4  
L6            462 S L5 AND L3  
L7            155 S L6 AND ELECTRODE#  
L8            432 S L6 AND (ELECTRODE# OR ANODE# OR CATHODE#)  
S 50-21-5/REG# OR 56-40-6/REG# OR 56-41-7/REG# OR 56-84-8/RE

FILE 'REGISTRY' ENTERED AT 14:59:41 ON 17 NOV 2008  
L9            1 S 110-17-8/RN

FILE 'CAPLUS' ENTERED AT 14:59:41 ON 17 NOV 2008  
L10          15717 S L9  
S 50-21-5/REG# OR 56-40-6/REG# OR 56-41-7/REG# OR 56-84-8/RE

L11 FILE 'REGISTRY' ENTERED AT 15:00:36 ON 17 NOV 2008  
1 S 56-87-1/RN

L12 FILE 'CAPLUS' ENTERED AT 15:00:37 ON 17 NOV 2008  
55791 S L11

L13 FILE 'REGISTRY' ENTERED AT 15:00:37 ON 17 NOV 2008  
1 S 56-86-0/RN

L14 FILE 'CAPLUS' ENTERED AT 15:00:37 ON 17 NOV 2008  
76293 S L13

L15 FILE 'REGISTRY' ENTERED AT 15:00:38 ON 17 NOV 2008  
1 S 56-84-8/RN

L16 FILE 'CAPLUS' ENTERED AT 15:00:38 ON 17 NOV 2008  
46908 S L15

L17 FILE 'REGISTRY' ENTERED AT 15:00:39 ON 17 NOV 2008  
1 S 56-41-7/RN

L18 FILE 'CAPLUS' ENTERED AT 15:00:39 ON 17 NOV 2008  
47918 S L17

L19 FILE 'REGISTRY' ENTERED AT 15:00:40 ON 17 NOV 2008  
1 S 56-40-6/RN

L20 FILE 'CAPLUS' ENTERED AT 15:00:40 ON 17 NOV 2008  
65792 S L19

L21 FILE 'REGISTRY' ENTERED AT 15:00:40 ON 17 NOV 2008  
1 S 50-21-5/RN

L22 FILE 'CAPLUS' ENTERED AT 15:00:41 ON 17 NOV 2008  
62717 S L21

L23 236074 S L22 OR L20 OR L18 OR L16 OR L14 OR L12  
S 61-90-5/REG# OR 70-26-8/REG# OR 74-79-3/REG# OR 77-92-9/RE

L24 FILE 'REGISTRY' ENTERED AT 15:01:17 ON 17 NOV 2008  
1 S 110-17-8/RN

L25 FILE 'CAPLUS' ENTERED AT 15:01:17 ON 17 NOV 2008  
15717 S L24

L26 FILE 'REGISTRY' ENTERED AT 15:01:17 ON 17 NOV 2008  
1 S 110-16-7/RN

L27 FILE 'CAPLUS' ENTERED AT 15:01:17 ON 17 NOV 2008  
16848 S L26

L28 FILE 'REGISTRY' ENTERED AT 15:01:18 ON 17 NOV 2008  
1 S 110-15-6/RN

L29 FILE 'CAPLUS' ENTERED AT 15:01:18 ON 17 NOV 2008  
31313 S L28

L30 FILE 'REGISTRY' ENTERED AT 15:01:19 ON 17 NOV 2008  
1 S 107-21-1/RN

L31 FILE 'CAPLUS' ENTERED AT 15:01:19 ON 17 NOV 2008  
54546 S L30

L32 FILE 'REGISTRY' ENTERED AT 15:01:20 ON 17 NOV 2008  
1 S 90-64-2/RN

L33 FILE 'CAPLUS' ENTERED AT 15:01:20 ON 17 NOV 2008  
4069 S L32

L34 FILE 'REGISTRY' ENTERED AT 15:01:20 ON 17 NOV 2008  
1 S 87-69-4/RN

L35 FILE 'CAPLUS' ENTERED AT 15:01:21 ON 17 NOV 2008  
24133 S L34

L36 FILE 'REGISTRY' ENTERED AT 15:01:21 ON 17 NOV 2008  
1 S 79-14-1/RN

L37 FILE 'CAPLUS' ENTERED AT 15:01:21 ON 17 NOV 2008  
10725 S L36

L38 FILE 'REGISTRY' ENTERED AT 15:01:22 ON 17 NOV 2008  
1 S 77-92-9/RN

L39 FILE 'CAPLUS' ENTERED AT 15:01:22 ON 17 NOV 2008  
73205 S L38

L40 FILE 'REGISTRY' ENTERED AT 15:01:23 ON 17 NOV 2008  
1 S 74-79-3/RN

L41 FILE 'CAPLUS' ENTERED AT 15:01:23 ON 17 NOV 2008  
51955 S L40

L42 FILE 'REGISTRY' ENTERED AT 15:01:23 ON 17 NOV 2008  
1 S 70-26-8/RN

L43 FILE 'CAPLUS' ENTERED AT 15:01:23 ON 17 NOV 2008  
8664 S L42

L44 FILE 'REGISTRY' ENTERED AT 15:01:24 ON 17 NOV 2008  
1 S 61-90-5/RN

L45 FILE 'CAPLUS' ENTERED AT 15:01:24 ON 17 NOV 2008  
40764 S L44

L46 FILE 'CAPLUS' ENTERED AT 15:01:24 ON 17 NOV 2008  
258073 S L45 OR L43 OR L41 OR L39 OR L37 OR L35 OR L33 OR L31 OR L29 O  
S 110-94-1/REG# OR 111-46-6/REG# OR 123-76-2/REG# OR 124-04-

L47 FILE 'REGISTRY' ENTERED AT 15:02:30 ON 17 NOV 2008  
1 S 499-12-7/RN

L48 FILE 'CAPLUS' ENTERED AT 15:02:30 ON 17 NOV 2008  
1047 S L47

L49 FILE 'REGISTRY' ENTERED AT 15:02:31 ON 17 NOV 2008  
1 S 498-23-7/RN

L50 FILE 'CAPLUS' ENTERED AT 15:02:31 ON 17 NOV 2008  
1043 S L49

L51 FILE 'REGISTRY' ENTERED AT 15:02:31 ON 17 NOV 2008  
1 S 298-12-4/RN

L52 FILE 'CAPLUS' ENTERED AT 15:02:32 ON 17 NOV 2008  
5548 S L51

L53 FILE 'REGISTRY' ENTERED AT 15:02:32 ON 17 NOV 2008  
1 S 144-62-7/RN

L54 FILE 'CAPLUS' ENTERED AT 15:02:32 ON 17 NOV 2008  
36542 S L53

L55 FILE 'REGISTRY' ENTERED AT 15:02:33 ON 17 NOV 2008  
1 S 141-82-2/RN

L56 FILE 'CAPLUS' ENTERED AT 15:02:33 ON 17 NOV 2008  
14876 S L55

L57 FILE 'REGISTRY' ENTERED AT 15:02:33 ON 17 NOV 2008  
1 S 127-17-3/RN

L58 FILE 'CAPLUS' ENTERED AT 15:02:34 ON 17 NOV 2008  
25465 S L57

L59 FILE 'REGISTRY' ENTERED AT 15:02:34 ON 17 NOV 2008  
1 S 124-04-9/RN

L60 FILE 'CAPLUS' ENTERED AT 15:02:34 ON 17 NOV 2008  
16236 S L59

L61 FILE 'REGISTRY' ENTERED AT 15:02:35 ON 17 NOV 2008  
1 S 123-76-2/RN

L62 FILE 'CAPLUS' ENTERED AT 15:02:35 ON 17 NOV 2008  
2735 S L61

L63 FILE 'REGISTRY' ENTERED AT 15:02:36 ON 17 NOV 2008  
1 S 111-46-6/RN

L64 FILE 'CAPLUS' ENTERED AT 15:02:36 ON 17 NOV 2008  
15270 S L63

L65 FILE 'REGISTRY' ENTERED AT 15:02:36 ON 17 NOV 2008  
1 S 110-94-1/RN

L66 FILE 'CAPLUS' ENTERED AT 15:02:36 ON 17 NOV 2008  
6189 S L65

L67 FILE 'CAPLUS' ENTERED AT 15:02:36 ON 17 NOV 2008  
108496 S L66 OR L64 OR L62 OR L60 OR L58 OR L56 OR L54 OR L52 OR L50 O  
S 6915-15-7/REG# OR 28854-76-4/REG# OR 35054-79-6/REG# OR 11

L68 FILE 'REGISTRY' ENTERED AT 15:03:27 ON 17 NOV 2008  
1 S 151677-68-8/RN

L69 FILE 'CAPLUS' ENTERED AT 15:03:27 ON 17 NOV 2008  
7 S L68

L70 FILE 'REGISTRY' ENTERED AT 15:03:28 ON 17 NOV 2008  
1 S 111937-70-3/RN

L71 FILE 'CAPLUS' ENTERED AT 15:03:28 ON 17 NOV 2008  
32 S L70

L72 FILE 'REGISTRY' ENTERED AT 15:03:28 ON 17 NOV 2008  
1 S 35054-79-6/RN

L73 FILE 'CAPLUS' ENTERED AT 15:03:29 ON 17 NOV 2008  
286 S L72

FILE 'REGISTRY' ENTERED AT 15:03:29 ON 17 NOV 2008  
L74 1 S 28854-76-4/RN

FILE 'CAPLUS' ENTERED AT 15:03:29 ON 17 NOV 2008  
L75 74 S L74

FILE 'REGISTRY' ENTERED AT 15:03:30 ON 17 NOV 2008  
L76 1 S 6915-15-7/RN

FILE 'CAPLUS' ENTERED AT 15:03:30 ON 17 NOV 2008  
L77 24029 S L76  
L78 24346 S L77 OR L75 OR L73 OR L71 OR L69

=> 16 and (19-178)

L6 IS NOT A RECOGNIZED COMMAND

The previous command name entered was not recognized by the system.  
For a list of commands available to you in the current file, enter  
"HELP COMMANDS" at an arrow prompt (>).

=> s l10 or l23 or l46 or l67 or l78  
L79 498661 L10 OR L23 OR L46 OR L67 OR L78

=> d his

(FILE 'HOME' ENTERED AT 14:46:54 ON 17 NOV 2008)

FILE 'CAPLUS' ENTERED AT 14:47:20 ON 17 NOV 2008  
E US20070152185/PN  
L1 1 S E3  
S 15365-14-7/REG# AND 7440-44-0/REG#

FILE 'REGISTRY' ENTERED AT 14:48:47 ON 17 NOV 2008  
L2 1 S 7440-44-0/RN

FILE 'CAPLUS' ENTERED AT 14:48:47 ON 17 NOV 2008  
L3 387474 S L2

FILE 'REGISTRY' ENTERED AT 14:48:47 ON 17 NOV 2008  
L4 1 S 15365-14-7/RN

FILE 'CAPLUS' ENTERED AT 14:48:48 ON 17 NOV 2008  
L5 1288 S L4  
L6 462 S L5 AND L3  
L7 155 S L6 AND ELECTRODE#  
L8 432 S L6 AND (ELECTRODE# OR ANODE# OR CATHODE#)  
S 50-21-5/REG# OR 56-40-6/REG# OR 56-41-7/REG# OR 56-84-8/RE

FILE 'REGISTRY' ENTERED AT 14:59:41 ON 17 NOV 2008  
L9 1 S 110-17-8/RN

FILE 'CAPLUS' ENTERED AT 14:59:41 ON 17 NOV 2008  
L10 15717 S L9  
S 50-21-5/REG# OR 56-40-6/REG# OR 56-41-7/REG# OR 56-84-8/RE

FILE 'REGISTRY' ENTERED AT 15:00:36 ON 17 NOV 2008  
L11 1 S 56-87-1/RN

FILE 'CAPLUS' ENTERED AT 15:00:37 ON 17 NOV 2008  
L12 55791 S L11

FILE 'REGISTRY' ENTERED AT 15:00:37 ON 17 NOV 2008  
L13 1 S 56-86-0/RN

FILE 'CAPLUS' ENTERED AT 15:00:37 ON 17 NOV 2008  
L14        76293 S L13

FILE 'REGISTRY' ENTERED AT 15:00:38 ON 17 NOV 2008  
L15        1 S 56-84-8/RN

FILE 'CAPLUS' ENTERED AT 15:00:38 ON 17 NOV 2008  
L16        46908 S L15

FILE 'REGISTRY' ENTERED AT 15:00:39 ON 17 NOV 2008  
L17        1 S 56-41-7/RN

FILE 'CAPLUS' ENTERED AT 15:00:39 ON 17 NOV 2008  
L18        47918 S L17

FILE 'REGISTRY' ENTERED AT 15:00:40 ON 17 NOV 2008  
L19        1 S 56-40-6/RN

FILE 'CAPLUS' ENTERED AT 15:00:40 ON 17 NOV 2008  
L20        65792 S L19

FILE 'REGISTRY' ENTERED AT 15:00:40 ON 17 NOV 2008  
L21        1 S 50-21-5/RN

FILE 'CAPLUS' ENTERED AT 15:00:41 ON 17 NOV 2008  
L22        62717 S L21

FILE 'CAPLUS' ENTERED AT 15:00:41 ON 17 NOV 2008  
L23        236074 S L22 OR L20 OR L18 OR L16 OR L14 OR L12  
            S 61-90-5/REG# OR 70-26-8/REG# OR 74-79-3/REG# OR 77-92-9/RE

FILE 'REGISTRY' ENTERED AT 15:01:17 ON 17 NOV 2008  
L24        1 S 110-17-8/RN

FILE 'CAPLUS' ENTERED AT 15:01:17 ON 17 NOV 2008  
L25        15717 S L24

FILE 'REGISTRY' ENTERED AT 15:01:17 ON 17 NOV 2008  
L26        1 S 110-16-7/RN

FILE 'CAPLUS' ENTERED AT 15:01:17 ON 17 NOV 2008  
L27        16848 S L26

FILE 'REGISTRY' ENTERED AT 15:01:18 ON 17 NOV 2008  
L28        1 S 110-15-6/RN

FILE 'CAPLUS' ENTERED AT 15:01:18 ON 17 NOV 2008  
L29        31313 S L28

FILE 'REGISTRY' ENTERED AT 15:01:19 ON 17 NOV 2008  
L30        1 S 107-21-1/RN

FILE 'CAPLUS' ENTERED AT 15:01:19 ON 17 NOV 2008  
L31        54546 S L30

FILE 'REGISTRY' ENTERED AT 15:01:20 ON 17 NOV 2008  
L32        1 S 90-64-2/RN

FILE 'CAPLUS' ENTERED AT 15:01:20 ON 17 NOV 2008  
L33        4069 S L32

FILE 'REGISTRY' ENTERED AT 15:01:20 ON 17 NOV 2008  
L34        1 S 87-69-4/RN

FILE 'CAPLUS' ENTERED AT 15:01:21 ON 17 NOV 2008  
L35        24133 S L34

FILE 'REGISTRY' ENTERED AT 15:01:21 ON 17 NOV 2008  
L36        1 S 79-14-1/RN

FILE 'CAPLUS' ENTERED AT 15:01:21 ON 17 NOV 2008  
L37        10725 S L36

FILE 'REGISTRY' ENTERED AT 15:01:22 ON 17 NOV 2008  
L38        1 S 77-92-9/RN

FILE 'CAPLUS' ENTERED AT 15:01:22 ON 17 NOV 2008  
L39        73205 S L38

FILE 'REGISTRY' ENTERED AT 15:01:23 ON 17 NOV 2008  
L40        1 S 74-79-3/RN

FILE 'CAPLUS' ENTERED AT 15:01:23 ON 17 NOV 2008  
L41        51955 S L40

FILE 'REGISTRY' ENTERED AT 15:01:23 ON 17 NOV 2008  
L42        1 S 70-26-8/RN

FILE 'CAPLUS' ENTERED AT 15:01:23 ON 17 NOV 2008  
L43        8664 S L42

FILE 'REGISTRY' ENTERED AT 15:01:24 ON 17 NOV 2008  
L44        1 S 61-90-5/RN

FILE 'CAPLUS' ENTERED AT 15:01:24 ON 17 NOV 2008  
L45        40764 S L44

L46        258073 S L45 OR L43 OR L41 OR L39 OR L37 OR L35 OR L33 OR L31 OR L29 O  
            S 110-94-1/REG# OR 111-46-6/REG# OR 123-76-2/REG# OR 124-04-

FILE 'REGISTRY' ENTERED AT 15:02:30 ON 17 NOV 2008  
L47        1 S 499-12-7/RN

FILE 'CAPLUS' ENTERED AT 15:02:30 ON 17 NOV 2008  
L48        1047 S L47

FILE 'REGISTRY' ENTERED AT 15:02:31 ON 17 NOV 2008  
L49        1 S 498-23-7/RN

FILE 'CAPLUS' ENTERED AT 15:02:31 ON 17 NOV 2008  
L50        1043 S L49

FILE 'REGISTRY' ENTERED AT 15:02:31 ON 17 NOV 2008  
L51        1 S 298-12-4/RN

FILE 'CAPLUS' ENTERED AT 15:02:32 ON 17 NOV 2008  
L52        5548 S L51

FILE 'REGISTRY' ENTERED AT 15:02:32 ON 17 NOV 2008  
L53        1 S 144-62-7/RN

FILE 'CAPLUS' ENTERED AT 15:02:32 ON 17 NOV 2008  
L54        36542 S L53

FILE 'REGISTRY' ENTERED AT 15:02:33 ON 17 NOV 2008  
L55        1 S 141-82-2/RN

FILE 'CAPLUS' ENTERED AT 15:02:33 ON 17 NOV 2008  
L56        14876 S L55

FILE 'REGISTRY' ENTERED AT 15:02:33 ON 17 NOV 2008  
L57        1 S 127-17-3/RN

FILE 'CAPLUS' ENTERED AT 15:02:34 ON 17 NOV 2008  
L58        25465 S L57

FILE 'REGISTRY' ENTERED AT 15:02:34 ON 17 NOV 2008  
L59        1 S 124-04-9/RN

FILE 'CAPLUS' ENTERED AT 15:02:34 ON 17 NOV 2008  
L60        16236 S L59

FILE 'REGISTRY' ENTERED AT 15:02:35 ON 17 NOV 2008  
L61        1 S 123-76-2/RN

FILE 'CAPLUS' ENTERED AT 15:02:35 ON 17 NOV 2008  
L62        2735 S L61

FILE 'REGISTRY' ENTERED AT 15:02:36 ON 17 NOV 2008  
L63        1 S 111-46-6/RN

FILE 'CAPLUS' ENTERED AT 15:02:36 ON 17 NOV 2008  
L64        15270 S L63

FILE 'REGISTRY' ENTERED AT 15:02:36 ON 17 NOV 2008  
L65        1 S 110-94-1/RN

FILE 'CAPLUS' ENTERED AT 15:02:36 ON 17 NOV 2008  
L66        6189 S L65

L67        108496 S L66 OR L64 OR L62 OR L60 OR L58 OR L56 OR L54 OR L52 OR L50 O  
            S 6915-15-7/REG# OR 28854-76-4/REG# OR 35054-79-6/REG# OR 11

FILE 'REGISTRY' ENTERED AT 15:03:27 ON 17 NOV 2008  
L68        1 S 151677-68-8/RN

FILE 'CAPLUS' ENTERED AT 15:03:27 ON 17 NOV 2008  
L69        7 S L68

FILE 'REGISTRY' ENTERED AT 15:03:28 ON 17 NOV 2008  
L70        1 S 111937-70-3/RN

FILE 'CAPLUS' ENTERED AT 15:03:28 ON 17 NOV 2008  
L71        32 S L70

FILE 'REGISTRY' ENTERED AT 15:03:28 ON 17 NOV 2008  
L72        1 S 35054-79-6/RN

FILE 'CAPLUS' ENTERED AT 15:03:29 ON 17 NOV 2008  
L73        286 S L72

FILE 'REGISTRY' ENTERED AT 15:03:29 ON 17 NOV 2008  
L74        1 S 28854-76-4/RN

FILE 'CAPLUS' ENTERED AT 15:03:29 ON 17 NOV 2008  
L75        74 S L74

FILE 'REGISTRY' ENTERED AT 15:03:30 ON 17 NOV 2008  
L76        1 S 6915-15-7/RN

FILE 'CAPLUS' ENTERED AT 15:03:30 ON 17 NOV 2008  
L77 24029 S L76  
L78 24346 S L77 OR L75 OR L73 OR L71 OR L69  
L79 498661 S L10 OR L23 OR L46 OR L67 OR L78

=> s 16 and 179  
L80 21 L6 AND L79

=> s 18 and 180  
L81 21 L8 AND L80

=> d 1-21 all

L81 ANSWER 1 OF 21 CAPLUS COPYRIGHT 2008 ACS on STN  
AN 2008:1169373 CAPLUS  
DN 149:474211  
ED Entered STN: 30 Sep 2008  
TI High-density olivine-type lithium ferrous phosphate as cathode material of lithium-ion battery and its preparation method  
IN Lu, Xiangyang; Zuo, Yicun  
PA Guangzhou Rongjie Material Science and Technology Co., Ltd., Peop. Rep. China  
SO Faming Zhuanli Shenqing Gongkai Shuomingshu, 8pp.  
CODEN: CNXXEV  
DT Patent  
LA Chinese  
CC 49-5 (Industrial Inorganic Chemicals)  
FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	CN 101269808	A	20080924	CN 2008-10026597	20080305
PRAI	CN 2008-10026597		20080305		

CLASS  
PATENT NO. CLASS PATENT FAMILY CLASSIFICATION CODES  
----- -----  
CN 101269808 IPCI C01B0025-45 [I,A]; C01B0025-00 [I,C\*]; H01M0004-58 [N,A]  
AB The title lithium ferrous phosphate material has olivine structure, average grain size of 1-8  $\mu\text{m}$ , apparent d.  $\geq 0.5 \text{ g/cm}^3$ , tap d. of 1.0-1.4  $\text{g/cm}^3$ , sp. surface area  $\leq 25 \text{ m}^2/\text{g}$ , and initial specific discharge capacity (1C)  $\geq 130 \text{ mAh/g}$ . The title method comprises dispersing Fe source, P source, Li source, doping element compound, and carbon source in dispersant, ball-milling for 1-12 h, drying under nonoxidative atmospheric

at 250-550° for 2-20 h, grinding, heating under reductive atmospheric at 15-40°/min, calcining at 500-900° for 3-15 h, and cooling at 15-40°/min to <40°. The mol. ratio of Fe source, P source, Li source and doping element is (0.7-1):1:(0.98-1.06):(0.02-0.3). The product has small particle size, narrow particle size distribution, high purity, and good electrochem. performance.

ST olivine lithium ferrous phosphate cathode lithium ion battery  
prepns

IT Density  
(apparent and tap; high-d. olivine-type lithium ferrous phosphate as cathode material of lithium-ion battery and its preparation method)

IT Ball milling  
Calcination  
Cathodes  
Crystal structure  
Electric properties  
Grinding (size reduction)

Particle size  
Solid phase synthesis  
(high-d. olivine-type lithium ferrous phosphate as cathode material of lithium-ion battery and its preparation method)

IT Secondary batteries  
(lithium; high-d. olivine-type lithium ferrous phosphate as cathode material of lithium-ion battery and its preparation method)

IT Surface area  
(specific; high-d. olivine-type lithium ferrous phosphate as cathode material of lithium-ion battery and its preparation method)

IT 7439-96-5, Manganese, uses  
RL: MOA (Modifier or additive use); USES (Uses)  
(dopant; high-d. olivine-type lithium ferrous phosphate as cathode material of lithium-ion battery and its preparation method)

IT 7440-44-0P, Carbon, preparation 15365-14-7P  
RL: IMF (Industrial manufacture); PRP (Properties); TEM (Technical or engineered material use); PREP (Preparation); USES (Uses)  
(high-d. olivine-type lithium ferrous phosphate as cathode material of lithium-ion battery and its preparation method)

IT 7439-95-4, Magnesium, uses 7439-98-7, Molybdenum, uses 7440-02-0, Nickel, uses  
RL: MOA (Modifier or additive use); USES (Uses)  
(high-d. olivine-type lithium ferrous phosphate as cathode material of lithium-ion battery and its preparation method)

IT 64-17-5, Ethanol, uses 1333-74-0, Hydrogen, uses 7727-37-9, Nitrogen, uses 7732-18-5, Water, uses  
RL: NUU (Other use, unclassified); USES (Uses)  
(high-d. olivine-type lithium ferrous phosphate as cathode material of lithium-ion battery and its preparation method)

IT 50-99-7, Glucose, reactions 57-50-1, Sucrose, reactions 77-92-9, Citric acid, reactions 546-89-4, Lithium acetate 554-13-2, Lithium carbonate 598-62-9, Manganese carbonate 1309-42-8, Magnesium hydroxide 1313-27-5, Molybdenum trioxide, reactions 1317-34-6, Manganese sesquioxide 7447-41-8, Lithium chloride, reactions 7664-38-2, Phosphoric acid, reactions 7705-08-0, Ferric chloride, reactions 7722-76-1, Ammonium dihydrogen phosphate 7783-28-0, Diammonium hydrogen phosphate 7790-69-4, Lithium nitrate 9003-05-8, Polyacrylamide 9003-07-0, Polypropylene 9004-53-9, Dextrin 9005-25-8, Starch, reactions 10045-86-0, Ferric phosphate 10138-04-2, Ammonium ferric sulfate 10377-60-3, Magnesium nitrate 10377-66-9, Manganous nitrate 10381-36-9, Nickelous phosphate 10421-48-4, Ferric nitrate 12057-24-8, Lithium oxide, reactions 14154-09-7, Manganese phosphate  
RL: RCT (Reactant); RACT (Reactant or reagent)  
(high-d. olivine-type lithium ferrous phosphate as cathode material of lithium-ion battery and its preparation method)

L81 ANSWER 2 OF 21 CAPLUS COPYRIGHT 2008 ACS on STN  
AN 2008:669636 CAPLUS  
DN 149:13781  
ED Entered STN: 05 Jun 2008  
TI Cathode active mass for secondary lithium batteries, and their manufacture, and the batteries  
IN Oshita, Itaru; Kanzaki, Kazuo  
PA Hitachi Maxell Ltd., Japan  
SO Jpn. Kokai Tokkyo Koho, 14pp.  
CODEN: JKXXAF  
DT Patent  
LA Japanese  
CC 52-2 (Electrochemical, Radiational, and Thermal Energy Technology)  
FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
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PI JP 2008130526 A 20080605 JP 2006-317924 20061127  
PRAI JP 2006-317924 20061127

## CLASS

PATENT NO.	CLASS	PATENT FAMILY CLASSIFICATION CODES
JP 2008130526	IPCI	H01M0004-58 [I,A]; H01M0004-02 [I,A]; H01M0010-40 [I,A]; H01M0010-36 [I,C*]; C01B0025-45 [I,A]; C01B0025-00 [I,C*]
	IPCR	H01M0004-58 [I,C]; H01M0004-58 [I,A]; C01B0025-00 [I,C]; C01B0025-45 [I,A]; H01M0004-02 [I,C]; H01M0010-36 [I,C]; H01M0010-40 [I,A]
	FTERM	5H029/AJ02; 5H029/AJ03; 5H029/AM12; 5H029/AM16; 5H029/CJ02; 5H029/CJ08; 5H029/CJ28; 5H029/DJ08; 5H029/DJ15; 5H029/DJ16; 5H029/EJ04; 5H029/HJ02; 5H029/HJ05; 5H029/HJ14; 5H050/AA02; 5H050/AA08; 5H050/BA17; 5H050/CA01; 5H050/DA02; 5H050/DA10; 5H050/EA01; 5H050/EA08; 5H050/EA09; 5H050/FA16; 5H050/FA17; 5H050/GA02; 5H050/GA10; 5H050/GA27; 5H050/HA02; 5H050/HA05; 5H050/HA14
AB		The active mass have olivine-type lithium iron phosphate primary particles and carbon-containing secondary particles, and the secondary particles have approx. spindle-, rhombus- or oval shape. The active mass is manufactured by a process including steps of (1) mixing lithium iron phosphate feedstock, carbonaceous materials, and C2-4 compds. bearing 2-3 hydroxy groups, and (2) heat treatment of the mixts. by hydrothermal crystallization, glycothermal process, or combination of two processes. Secondary Li batteries employing the cathode active mass are capable of high-speed charging and discharging and show high discharge capacity.
ST		battery cathode lithium iron phosphate composite carbon; hydrothermal crystn lithium iron phosphate composite battery cathode; glycothermal process lithium iron phosphate composite battery cathode
IT	Carbon black, uses	RL: TEM (Technical or engineered material use); USES (Uses) (acetylene black, ketjen black, composites with lithium iron phosphates, cathode active mass; manufacture of lithium iron phosphate-carbon composite granules as secondary Li battery cathodes)
IT	Nanotubes	(carbon, composites with lithium iron phosphates, cathode active mass; manufacture of lithium iron phosphate-carbon composite granules as secondary Li battery cathodes)
IT	Carbon fibers, uses	Fullerenes RL: TEM (Technical or engineered material use); USES (Uses) (composites with lithium iron phosphates, cathode active mass; manufacture of lithium iron phosphate-carbon composite granules as secondary Li battery cathodes)
IT	Secondary batteries	(lithium; manufacture of lithium iron phosphate-carbon composite granules as secondary Li battery cathodes)
IT	Battery cathodes	Hydrothermal crystallization (manufacture of lithium iron phosphate-carbon composite granules as secondary Li battery cathodes)
IT	Crystallization	(thermal, solvothermal; manufacture of lithium iron phosphate-carbon composite granules as secondary Li battery cathodes)
IT	7440-44-0, Carbon, uses	7782-42-5, Graphite, uses RL: TEM (Technical or engineered material use); USES (Uses) (composites with lithium iron phosphates, cathode active mass; manufacture of lithium iron phosphate-carbon composite granules as

secondary Li battery cathodes)

IT 56-81-5, Glycerol, uses 57-55-6, Propylene glycol, uses 107-21-1  
, Ethylene glycol, uses 107-88-0, 1,3-Butanediol 110-63-4,  
1,4-Butanediol, uses 504-63-2, 1,3-Propanediol 3068-00-6,  
1,2,4-Butanetriol 4435-50-1, 1,2,3-Butanetriol  
RL: NUU (Other use, unclassified); USES (Uses)  
(granulating agents; in manufacture of lithium iron phosphate-carbon composite granules as secondary Li battery cathodes)

IT 1310-65-2, Lithium hydroxide 1310-66-3, Lithium hydroxide monohydrate 7664-38-2, Phosphoric acid, uses 7720-78-7 7758-94-3, Ferrous chloride 7782-63-0, Iron sulfate heptahydrate 10045-89-3 13463-43-9, Ferrous sulfate hydrate 23838-02-0, Ferrous chloride hydrate 62586-14-5, Ammonium iron sulfate hydrate  
RL: TEM (Technical or engineered material use); USES (Uses)  
(in manufacture of lithium iron phosphate-carbon composite granules as secondary Li battery cathodes)

IT 15365-14-7P, Iron lithium phosphate (LiFePO<sub>4</sub>)  
RL: IMF (Industrial manufacture); TEM (Technical or engineered material use); PREP (Preparation); USES (Uses)  
(olivine-type, composites with carbon, cathode active mass;  
manufacture of lithium iron phosphate-carbon composite granules as secondary Li battery cathodes)

L81 ANSWER 3 OF 21 CAPLUS COPYRIGHT 2008 ACS on STN  
AN 2008:595199 CAPLUS  
DN 149:13704  
ED Entered STN: 19 May 2008  
TI Production of LiFePO<sub>4</sub>/C cathode material for lithium-ion batteries  
IN Liu, Xinbao; Jia, Xiaolin; Ma, Meipin; Liu, Yufei; Zhou, Yonggang; Wu, Jiangfeng  
PA Zhengzhou Dlg Battery Co., Ltd., Peop. Rep. China  
SO Faming Zhuanli Shenqing Gongkai Shuomingshu, 16pp.  
CODEN: CNXXEV  
DT Patent  
LA Chinese  
CC 52-2 (Electrochemical, Radiational, and Thermal Energy Technology)

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	CN 101179124	A	20080514	CN 2007-10055033	20070824
PRAI	CN 2007-10055033		20070824		

CLASS

PATENT NO.	CLASS	PATENT FAMILY CLASSIFICATION CODES
CN 101179124	IPCI	H01M0004-04 [I,A]; H01M0004-58 [I,A]; C01B0025-45 [I,A]; C01B0025-00 [I,C*]
	IPCR	H01M0004-04 [I,C]; H01M0004-04 [I,A]

AB This method entails: (a) preparing a solution of a Li compound, Fe compound, P compound and reductant at a molar ratio of 1:(0.8-1.2):(0.8-1.2):(0.3-1.0), adding Fe compound and Li compound to reductant, then adding P compound, mixing evenly under stirring, and aging, (b) drying under nonoxidative atmospheric to obtain dry gel, and grinding to obtain precursor, and (c) adding C compound 1.5-11.5% to the precursor, grinding, adding to a crucible, and microwave heating under nonoxidative atmospheric to obtain the final product. The method uses Cr<sup>3+</sup> and Co<sup>3+</sup> to modify LiFePO<sub>4</sub>/C. The method has the advantages of a decreased treatment time, enhanced yield, reduced cost and energy consumption and simplified process. The cathode material has the advantages of enhanced conductivity of LiFePO<sub>4</sub> and enhanced charge-discharge capacity and cycling ability.

ST iron lithium phosphate carbon cathode material lithium battery

IT Carbon black, reactions  
 RL: RCT (Reactant); RACT (Reactant or reagent)  
 (in production of LiFePO<sub>4</sub>/C cathode material for lithium-ion batteries)

IT Secondary batteries  
 (lithium; production of LiFePO<sub>4</sub>/C cathode material for lithium-ion batteries)

IT Battery cathodes  
 (production of LiFePO<sub>4</sub>/C cathode material for lithium-ion batteries)

IT 5931-89-5, Cobalt acetate 10103-47-6, Chromium nitrate 10124-43-3, Cobalt sulfate 10141-05-6, Cobalt nitrate 12336-95-7, Basic chromium sulfate 14489-25-9, Chromium sulfate 17593-70-3, Chromium acetate RL: MOA (Modifier or additive use); USES (Uses)  
 (in production of LiFePO<sub>4</sub>/C cathode material for lithium-ion batteries)

IT 7440-37-1, Argon, uses 7440-59-7, Helium, uses 7727-37-9, Nitrogen, uses  
 RL: NUU (Other use, unclassified); USES (Uses)  
 (in production of LiFePO<sub>4</sub>/C cathode material for lithium-ion batteries)

IT 50-99-7, Glucose, reactions 57-50-1, Sucrose, reactions 63-42-3, Lactose 69-79-4, Maltose 546-89-4, Lithium acetate 1310-65-2, Lithium hydroxide 7664-38-2, Phosphoric acid, reactions 7782-42-5, Graphite, reactions 7783-28-0, Diammonium hydrogen phosphate 7790-69-4, Lithium nitrate 10028-22-5, Ferric sulfate 10124-31-9, Ammonium phosphate 10377-52-3, Lithium phosphate 10421-48-4, Ferric nitrate  
 RL: RCT (Reactant); RACT (Reactant or reagent)  
 (in production of LiFePO<sub>4</sub>/C cathode material for lithium-ion batteries)

IT 50-81-7, Ascorbic acid, reactions 79-14-1, Glycolic acid, reactions  
 RL: RGT (Reagent); RACT (Reactant or reagent)  
 (in production of LiFePO<sub>4</sub>/C cathode material for lithium-ion batteries)

IT 7440-44-0P, Carbon, uses 15365-14-7P, Iron lithium phosphate (FeLiPO<sub>4</sub>)  
 RL: IMF (Industrial manufacture); TEM (Technical or engineered material use); PREP (Preparation); USES (Uses)  
 (production of LiFePO<sub>4</sub>/C cathode material for lithium-ion batteries)

L81 ANSWER 4 OF 21 CAPLUS COPYRIGHT 2008 ACS on STN  
 AN 2007:1309501 CAPLUS  
 DN 147:525392  
 ED Entered STN: 16 Nov 2007  
 TI Cathode material for rechargeable batteries  
 IN Yang, Chih-Wei; Liu, Wen-Ren  
 PA Aquire Energy Co. Ltd., Taiwan  
 SO Eur. Pat. Appl., 19pp.  
 CODEN: EPXXDW

DT Patent  
 LA English  
 CC 52-2 (Electrochemical, Radiational, and Thermal Energy Technology)  
 FAN.CNT 8

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
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PI	EP 1855334	A2	20071114	EP 2007-251938	20070510
	R: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LI, LT, LU, LV, MC, MT, NL, PL, PT, RO, SE, SI, SK, TR, AL, BA, HR, MK, YU				

AT 385999	T	20080315	AT 2005-256174	20051003
CA 2588494	A1	20071111	CA 2007-2588494	20070510
KR 2007109903	A	20071115	KR 2007-45322	20070510
JP 2007305585	A	20071122	JP 2007-125731	20070510
IN 2007KO00730	A	20071123	IN 2007-KO730	20070510
CN 101304083	A	20081112	CN 2007-10103278	20070510
PRAI CN 2006-10080365	A	20060511		
EP 2005-256174	A	20051003		
CLASS				
PATENT NO.	CLASS	PATENT FAMILY CLASSIFICATION CODES		
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EP 1855334	IPCI	H01M0004-48 [I,A]; H01M0004-58 [I,A]; H01M0010-40 [I,A]; H01M0010-36 [I,C*]		
	IPCR	H01M0004-48 [I,C]; H01M0004-48 [I,A]; H01M0004-58 [I,C]; H01M0004-58 [I,A]; H01M0010-36 [I,C]; H01M0010-40 [I,A]		
AT 385999	IPCI	C01G0049-00 [I,C]; C01G0049-00 [I,A]; C01B0013-14 [I,C]; C01B0013-14 [I,A]; C01B0025-00 [I,C]; C01B0025-45 [I,A]; C01D0001-00 [I,C]; C01D0001-02 [I,A]; H01M0004-48 [I,C]; H01M0004-48 [I,A]; H01M0004-58 [I,C]; H01M0004-58 [I,A]		
	IPCR	C01G0049-00 [I,C]; C01G0049-00 [I,A]; C01B0013-14 [I,C]; C01B0013-14 [I,A]; C01B0025-00 [I,C]; C01B0025-45 [I,A]; C01D0001-00 [I,C]; C01D0001-02 [I,A]; H01M0004-48 [I,C]; H01M0004-48 [I,A]; H01M0004-58 [I,C]; H01M0004-58 [I,A]		
	ECLA	C01B025/37; H01M004/58D; M01P; M01P; M01P; T01M; T01M		
CA 2588494	IPCI	H01M0004-36 [I,A]; H01M0004-52 [I,A]; H01M0004-58 [I,A]; H01M0010-00 [I,A]		
	IPCR	H01M0004-58 [I,C]; H01M0004-58 [I,A]; H01M0004-36 [I,C]; H01M0004-36 [I,A]; H01M0004-52 [I,C]; H01M0004-52 [I,A]; H01M0010-00 [I,C]; H01M0010-00 [I,A]		
KR 2007109903	IPCI	H01M0004-58 [I,A]; H01M0004-62 [I,A]; H01M0010-02 [I,A]		
JP 2007305585	IPCI	H01M0004-58 [I,A]; H01M0004-36 [I,A]; H01M0004-62 [I,A]		
	IPCR	H01M0004-58 [I,C]; H01M0004-58 [I,A]; H01M0004-36 [I,C]; H01M0004-36 [I,A]; H01M0004-62 [I,C]; H01M0004-62 [I,A]		
	FTERM	5H050/AA02; 5H050/AA08; 5H050/AA12; 5H050/BA16; 5H050/CA01; 5H050/CB12; 5H050/DA10; 5H050/EA08; 5H050/EA12; 5H050/FA17; 5H050/FA18; 5H050/HA01; 5H050/HA02		
IN 2007KO00730	IPCI	H01M0004-58 [ICM,7]; H01M0010-02 [ICS,7]		
CN 101304083	IPCI	H01M0004-02 [I,A]; H01M0010-04 [I,A]		
AB	This cathode material consists of a compound with an olivine or NASICON structure and a conductive metal oxide. The rechargeable battery has an anode, an electrolyte and a cathode as described above.			
ST	cathode rechargeable battery			
IT	Battery cathodes Secondary batteries (cathode material for rechargeable batteries)			
IT	Oxides (inorganic), uses Phosphates, uses RL: TEM (Technical or engineered material use); USES (Uses) (cathode material for rechargeable batteries)			
IT	1314-13-2P, Zinc oxide (ZnO), uses 1344-28-1P, Alumina, uses 1344-70-3P, Copper oxide 7631-86-9P, Silica, uses 11099-11-9P, Vanadium oxide 11104-61-3P, Cobalt oxide 11129-60-5P, Manganese oxide 13463-67-7P, Titanium oxide (TiO2), uses 15365-14-7P, Iron lithium phosphate (FeLiPO4) 37275-76-6P, Aluminum zinc oxide RL: SPN (Synthetic preparation); TEM (Technical or engineered material			

use); PREP (Preparation); USES (Uses)  
(cathode material for rechargeable batteries)

IT 1307-96-6P, Cobaltous oxide, uses 1309-48-4P, Magnesium oxide (MgO),  
uses 1313-99-1P, Nickel oxide (NiO), uses 1317-38-0P, Cupric oxide,  
uses 1344-43-0P, Manganous oxide, uses  
RL: SPN (Synthetic preparation); TEM (Technical or engineered material  
use); PREP (Preparation); USES (Uses)  
(in cathode material for rechargeable batteries)

IT 1314-62-1, Vanadium oxide (V2O5), uses 7440-44-0, Carbon, uses  
RL: TEM (Technical or engineered material use); USES (Uses)  
(in cathode material for rechargeable batteries)

IT 57-50-1, Sucrose, reactions 77-92-9, Citric acid, reactions  
1310-65-2, Lithium hydroxide (Li(OH)) 3251-23-8 7439-89-6, Iron,  
reactions 7646-85-7, Zinc chloride (ZnCl2), reactions 7664-38-2,  
Phosphoric acid, reactions 10026-22-9, Cobalt nitrate (Co(NO3)2)  
hexahydrate 13478-00-7, Nickel nitrate (Ni(NO3)2) hexahydrate  
17141-63-8, Manganese nitrate (Mn(NO3)2) hexahydrate  
RL: RCT (Reactant); RACT (Reactant or reagent)  
(in preparation of cathode material for rechargeable batteries)

L81 ANSWER 5 OF 21 CAPLUS COPYRIGHT 2008 ACS on STN  
AN 2007:1292448 CAPLUS  
DN 148:148282  
ED Entered STN: 14 Nov 2007  
TI Effects of Reductive Conditions on the Microstructure and Electrochemical Properties of Sol-Gel Derived LiFePO4/C  
AU Lin, Yan; Pan, Hongge; Gao, Mingxia; Liu, Yongfeng  
CS Department of Materials Science and Engineering, Zhejiang University, Hangzhou, 310027, Peop. Rep. China  
SO Journal of the Electrochemical Society (2007), 154(12), A1124-A1128  
CODEN: JESOAN; ISSN: 0013-4651  
PB Electrochemical Society  
DT Journal  
LA English  
CC 52-2 (Electrochemical, Radiational, and Thermal Energy Technology)  
Section cross-reference(s): 49, 72  
AB A sol-gel method was used to prepare LiFePO4/C composites by using ferric iron as iron precursor. Effects of reductive conditions including the amts. of organic additive of citric acid and the H2 content in the sintering atmospheric of N2 on the microstructural characteristics and electrochem. properties of LiFePO4/C are studied. The LiFePO4/C sample prepared with citric acid and cations in a ratio of 1:2 under sintering atmospheric of 10% H2 + N2 shows a highest discharge capacity of 135 mAh/g at the rate of 0.1 C (1 C = 170 mA/g). A carbon content of .apprx.4% provides a satisfactory discharge capacity of LiFePO4. Citric acid can act as a reducing reagent; hydrogen at 10-20 volume% in the sintering atmospheric can also reduce Fe3+ to Fe2+.  
Fe2+ completely and a further suitable amount of electro-conductive Fe2P phase could possibly form. Iron phosphides play an important role in the improvement of the high rate capacity of LiFePO4/C and the reaction kinetics of lithium ion. At discharge rates >1 C, suitably increasing the amount of Fe2P to a range of 6-11% in the LiFePO4/C composite favors the rate performance of LiFePO4/C.  
ST reductive atm microstructure electrochem behavior iron lithium phosphate; secondary lithium battery cathode iron lithium phosphate discharge capacity  
IT Reduction  
(after sintering; effects of reductive conditions on microstructure and electrochem. properties of Sol-gel derived LiFePO4/C)  
IT Electric capacitance  
(discharge capacity; effects of reductive conditions on microstructure

and electrochem. properties of Sol-gel derived LiFePO<sub>4</sub>/C)

IT Controlled atmospheres  
(during sintering; effects of reductive conditions on microstructure and electrochem. properties of Sol-gel derived LiFePO<sub>4</sub>/C)

IT Battery cathodes  
Composites  
Cyclic voltammetry  
Ionic conductivity  
Sintering  
Sol-gel processing  
(effects of reductive conditions on microstructure and electrochem. properties of Sol-gel derived LiFePO<sub>4</sub>/C)

IT Carbon black, uses  
Fluoropolymers, uses  
RL: TEM (Technical or engineered material use); USES (Uses)  
(effects of reductive conditions on microstructure and electrochem. properties of Sol-gel derived LiFePO<sub>4</sub>/C)

IT Secondary batteries  
(lithium; effects of reductive conditions on microstructure and electrochem. properties of Sol-gel derived LiFePO<sub>4</sub>/C)

IT Microstructure  
(of cathode material; effects of reductive conditions on microstructure and electrochem. properties of Sol-gel derived LiFePO<sub>4</sub>/C)

IT Electric current-potential relationship  
(of cathodes during cyclic voltammetry; effects of reductive conditions on microstructure and electrochem. properties of Sol-gel derived LiFePO<sub>4</sub>/C)

IT 7440-44-0, Super P, uses  
RL: ANT (Analyte); TEM (Technical or engineered material use); ANST (Analytical study); USES (Uses)  
(activated; effects of reductive conditions on microstructure and electrochem. properties of Sol-gel derived LiFePO<sub>4</sub>/C)

IT 1309-37-1, Iron oxide (Fe<sub>2</sub>O<sub>3</sub>), analysis 36058-25-0, Iron lithium phosphate (Fe<sub>2</sub>Li<sub>3</sub>PO<sub>4</sub>)<sub>3</sub>  
RL: ANT (Analyte); FMU (Formation, unclassified); ANST (Analytical study); FORM (Formation, nonpreparative)  
(after sintering; effects of reductive conditions on microstructure and electrochem. properties of Sol-gel derived LiFePO<sub>4</sub>/C)

IT 15365-14-7P, Iron lithium phosphate (FeLiPO<sub>4</sub>)  
RL: ANT (Analyte); PRP (Properties); SPN (Synthetic preparation); TEM (Technical or engineered material use); ANST (Analytical study); PREP (Preparation); USES (Uses)  
(effects of reductive conditions on microstructure and electrochem. properties of Sol-gel derived LiFePO<sub>4</sub>/C)

IT 77-92-9, Citric acid, reactions 1333-74-0, Hydrogen, reactions 7722-76-1, Ammonium dihydrogen phosphate 7790-69-4, Lithium nitrate  
RL: RCT (Reactant); RACT (Reactant or reagent)  
(effects of reductive conditions on microstructure and electrochem. properties of Sol-gel derived LiFePO<sub>4</sub>/C)

IT 96-49-1, Ethylene carbonate 616-38-6, Dimethyl carbonate 7429-90-5, Aluminum, uses 7439-93-2, Lithium, uses 21324-40-3, Lithium hexafluorophosphate 24937-79-9, PVDF  
RL: TEM (Technical or engineered material use); USES (Uses)  
(effects of reductive conditions on microstructure and electrochem. properties of Sol-gel derived LiFePO<sub>4</sub>/C)

IT 1310-43-6, Iron phosphide (Fe<sub>2</sub>P) 26508-33-8, Iron phosphide (FeP)  
RL: ANT (Analyte); FMU (Formation, unclassified); ANST (Analytical study); FORM (Formation, nonpreparative)  
(formed in cathode; effects of reductive conditions on microstructure and electrochem. properties of Sol-gel derived LiFePO<sub>4</sub>/C)

IT 10377-52-3, Lithium phosphate (Li<sub>3</sub>PO<sub>4</sub>)  
 RL: FMU (Formation, unclassified); FORM (Formation, nonpreparative)  
 (formed in cathode; effects of reductive conditions on  
 microstructure and electrochem. properties of Sol-gel derived  
 LiFePO<sub>4</sub>/C)  
 IT 10421-48-4, Ferric nitrate  
 RL: RCT (Reactant); RACT (Reactant or reagent)  
 (precursor; effects of reductive conditions on microstructure and  
 electrochem. properties of Sol-gel derived LiFePO<sub>4</sub>/C)

RE.CNT 17 THERE ARE 17 CITED REFERENCES AVAILABLE FOR THIS RECORD  
 RE

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 Electrochemical Society of Japan Meeting Abstracts 1999, V99-2
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- (15) Xu, Y; J Power Sources 2006, V160, P570 CAPLUS
- (16) Yang, J; Electrochem Solid-State Lett 2004, V7, PA515 CAPLUS
- (17) Yang, J; J Electrochem Soc 2006, V153, PA716 CAPLUS

L81 ANSWER 6 OF 21 CAPLUS COPYRIGHT 2008 ACS on STN

AN 2007:1237548 CAPLUS

DN 147:505381

ED Entered STN: 01 Nov 2007

TI Cathode material for manufacturing a rechargeable battery

IN Yang, Chih-Wei

PA Aquire Energy Co. Ltd., Taiwan

SO Eur. Pat. Appl., 21pp.

CODEN: EPXXDW

DT Patent

LA English

CC 52-2 (Electrochemical, Radiational, and Thermal Energy Technology)  
 Section cross-reference(s): 49

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	EP 1850409	A1	20071031	EP 2007-251680	20070423
	R: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LI, LT, LU, LV, MC, MT, NL, PL, PT, RO, SE, SI, SK, TR, AL, BA, HR, MK, YU				
	CN 101064367	A	20071031	CN 2006-10074964	20060425
	CA 2585594	A1	20071025	CA 2007-2585594	20070420
	KR 2007105266	A	20071030	KR 2007-39924	20070424
	KR 809570	B1	20080304		
	JP 2007294461	A	20071108	JP 2007-114024	20070424
	IN 2007KO00638	A	20080725	IN 2007-KO638	20070425
PRAI	CN 2006-10074964	A	20060425		

CLASS

PATENT NO.	CLASS	PATENT FAMILY CLASSIFICATION CODES
EP 1850409	IPCI	H01M0004-58 [I,A]; H01M0010-40 [I,A]; H01M0010-36 [I,C*]

CN	101064367	IPCI	H01M0004-02 [I,A]; H01M0004-48 [I,A]; H01M0004-58 [I,A]; C01B0025-45 [I,A]; C01B0025-00 [I,C*]; H01M0010-40 [I,A]; H01M0010-36 [I,C*]
		IPCR	H01M0004-02 [I,C]; H01M0004-02 [I,A]
CA	2585594	IPCI	H01M0004-58 [I,A]; H01M0010-02 [I,A]
		IPCR	H01M0004-58 [I,C]; H01M0004-58 [I,A]; H01M0010-02 [I,C]; H01M0010-02 [I,A]
KR	2007105266	IPCI	H01M0004-58 [I,A]; H01M0004-38 [I,A]
JP	2007294461	IPCI	H01M0004-58 [I,A]; H01M0004-36 [I,A]; H01M0004-02 [I,A]; H01M0004-62 [I,A]
		IPCR	H01M0004-58 [I,C]; H01M0004-58 [I,A]; H01M0004-02 [I,C]; H01M0004-02 [I,A]; H01M0004-36 [I,C]; H01M0004-36 [I,A]; H01M0004-62 [I,C]; H01M0004-62 [I,A]
		FTERM	5H050/AA02; 5H050/AA08; 5H050/BA17; 5H050/CA01; 5H050/CB07; 5H050/DA10; 5H050/DA11; 5H050/DA14; 5H050/EA08; 5H050/EA23; 5H050/EA24; 5H050/FA17; 5H050/FA18; 5H050/HA02; 5H050/HA05; 5H050/HA07
IN	2007K000638	IPCI	H01M0006-12 [ICM,7]; H01M0006-04 [ICM,7,C*]; H01M0006-46 [ICS,7]; H01M0006-42 [ICS,7,C*]

AB A cathode material includes crystalline nanometer-sized primary particles of a metal compound having one of olivine and NASICON structures and a particle size ranging from 10 to 500 nm, and micrometer-sized secondary particles having a particle size ranging from 1 to 50  $\mu\text{m}$ . Each of the micrometer-sized secondary particles is composed of the crystalline nanometer-sized primary particles.

ST cathode material rechargeable battery fabrication

IT Battery cathodes

Nanoparticles

Particle size

Secondary batteries

Surface area

(cathode material for manufacturing rechargeable battery)

IT Carbonaceous materials (technological products)

Fluoropolymers, uses

Styrene-butadiene rubber, uses

RL: MOA (Modifier or additive use); USES (Uses)

(cathode material for manufacturing rechargeable battery)

IT 50-99-7, Glucose, uses 57-50-1, Sucrose, uses 872-50-4, N-Methylpyrrolidone, uses 7446-70-0, Aluminum chloride (AlCl<sub>3</sub>), uses 7786-30-3, Magnesium chloride (MgCl<sub>2</sub>), uses 9000-11-7, CMC 24937-79-9, PVDF

RL: MOA (Modifier or additive use); USES (Uses)

(cathode material for manufacturing rechargeable battery)

IT 77-92-9, Citric acid, reactions 144-62-7, Oxalic acid, reactions 7439-89-6, Iron, reactions 7705-08-0, Ferric chloride, reactions 10421-48-4, Ferric nitrate

RL: RCT (Reactant); RACT (Reactant or reagent)

(cathode material for manufacturing rechargeable battery)

IT 15365-14-7P, Iron lithium phosphate felipo4 928163-03-5P

RL: SPN (Synthetic preparation); TEM (Technical or engineered material use); PREP (Preparation); USES (Uses)

(cathode material for manufacturing rechargeable battery)

IT 7440-44-0, Carbon, uses

RL: MOA (Modifier or additive use); USES (Uses)

(particles; cathode material for manufacturing rechargeable battery)

IT 9003-55-8

RL: MOA (Modifier or additive use); USES (Uses)

(styrene-butadiene rubber; cathode material for manufacturing rechargeable battery)

RE.CNT 3 THERE ARE 3 CITED REFERENCES AVAILABLE FOR THIS RECORD

RE

- (1) Aquire Energy Co Ltd; EP 1790617 A 2007 CAPLUS  
 (2) Sony Corp; EP 1094533 A 2001 CAPLUS  
 (3) Sony Corp; EP 1193783 A 2002 CAPLUS

L81 ANSWER 7 OF 21 CAPLUS COPYRIGHT 2008 ACS on STN  
 AN 2007:971184 CAPLUS  
 DN 147:280968  
 ED Entered STN: 31 Aug 2007  
 TI Manufacture of iron lithium phosphate electrode materials and secondary lithium batteries using them  
 IN Inamasu, Tokuo; Fujii, Akihiro; Nukuta, Toshiyuki  
 PA Gs Yuasa Corp., Japan  
 SO Jpn. Kokai Tokkyo Koho, 8pp.  
 CODEN: JKXXAF  
 DT Patent  
 LA Japanese  
 CC 52-2 (Electrochemical, Radiational, and Thermal Energy Technology)  
 FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI JP 2007220390	A	20070830	JP 2006-37533	20060215
PRAI JP 2006-37533		20060215		

CLASS

PATENT NO.	CLASS	PATENT FAMILY CLASSIFICATION CODES
JP 2007220390	IPCI	H01M0004-58 [I,A]; H01M0010-40 [N,A]; H01M0010-36 [N,C*]
	IPCR	H01M0004-58 [I,C]; H01M0004-58 [I,A]; H01M0010-36 [N,C]; H01M0010-40 [N,A]
	FTERM	5H029/AJ02; 5H029/AJ03; 5H029/AK03; 5H029/AL12; 5H029/AM03; 5H029/AM05; 5H029/AM07; 5H029/BJ12; 5H029/CJ02; 5H029/CJ08; 5H029/CJ28; 5H029/DJ16; 5H029/EJ01; 5H029/EJ04; 5H029/EJ12; 5H029/EJ14; 5H029/HJ14; 5H050/AA02; 5H050/AA08; 5H050/BA17; 5H050/CA07; 5H050/CA08; 5H050/CB12; 5H050/EA02; 5H050/EA08; 5H050/EA09; 5H050/EA10; 5H050/EA23; 5H050/EA24; 5H050/FA17; 5H050/GA02; 5H050/GA10; 5H050/GA27; 5H050/HA14; 5H050/HA20

AB The materials are manufactured by treatment of aqueous solns. containing Fe compds. and

conductive materials (e.g. carbon powder) with oxalic acid, and firing of the resulting Fe(II) oxalate with Li compds. and phosphoric acids. The materials and batteries show high utilization efficiency of iron lithium phosphate.

ST iron lithium phosphate lithium battery cathode; lithium battery cathode iron oxalate firing; battery cathode iron lithium phosphate carbon

IT Secondary batteries  
 (lithium; manufacture of LiFePO<sub>4</sub> cathodes containing carbon conductive materials for secondary lithium batteries from Fe compds. via Fe oxalate)

IT Battery cathodes  
 (manufacture of LiFePO<sub>4</sub> cathodes containing carbon conductive materials for secondary lithium batteries from Fe compds. via Fe oxalate)

IT Carbon black, uses  
 RL: TEM (Technical or engineered material use); USES (Uses)  
 (manufacture of LiFePO<sub>4</sub> cathodes containing carbon conductive materials for secondary lithium batteries from Fe compds. via Fe oxalate)

IT 516-03-0P, Iron (II) oxalate  
 RL: IMF (Industrial manufacture); PEP (Physical, engineering or chemical

process); PREP (Preparation); PROC (Process)  
(manufacture of LiFePO<sub>4</sub> cathodes containing carbon conductive materials for secondary lithium batteries from Fe compds. via Fe oxalate)

IT 15365-14-7P, Iron lithium phosphate (FeLiPO<sub>4</sub>)  
RL: IMF (Industrial manufacture); TEM (Technical or engineered material use); PREP (Preparation); USES (Uses)  
(manufacture of LiFePO<sub>4</sub> cathodes containing carbon conductive materials for secondary lithium batteries from Fe compds. via Fe oxalate)

IT 144-62-7, Oxalic acid, processes 554-13-2, Lithium carbonate  
7720-78-7, Iron(II) sulfate 7722-76-1, Ammonium dihydrogen phosphate  
RL: PEP (Physical, engineering or chemical process); PROC (Process)  
(manufacture of LiFePO<sub>4</sub> cathodes containing carbon conductive materials for secondary lithium batteries from Fe compds. via Fe oxalate)

IT 7440-44-0, Carbon, uses  
RL: TEM (Technical or engineered material use); USES (Uses)  
(manufacture of LiFePO<sub>4</sub> cathodes containing carbon conductive materials for secondary lithium batteries from Fe compds. via Fe oxalate)

L81 ANSWER 8 OF 21 CAPLUS COPYRIGHT 2008 ACS on STN  
AN 2007:470396 CAPLUS  
DN 148:154036  
ED Entered STN: 30 Apr 2007  
TI Improvement of the electrochemical performance of LiFePO<sub>4</sub> cathode composite material using a in situ pyrolysis carbon synthesis procedure  
AU Luo, Shaohua; Tang, Zilong; Lu, Junbiao; Li, Junrong; Zhang, Zhongtai  
CS State Key Laboratory of New Ceramics and Fine Processing, Department of Materials Science and Engineering, Tsinghua University, Beijing, 100084, Peop. Rep. China  
SO Key Engineering Materials (2007), 336-338(Pt. 1, High-Performance Ceramics IV), 466-469  
CODEN: KEMAEY; ISSN: 1013-9826  
PB Trans Tech Publications Ltd.  
DT Journal  
LA English  
CC 72-2 (Electrochemistry)  
Section cross-reference(s): 52  
AB LiFePO<sub>4</sub>/C composite cathode material was prepared by pelletting and subsequent pyrolytic cracking process in N<sub>2</sub> atmosphere with C source of polyvinyl alc. (PVA). XRD crystal anal. indicates that single LiFePO<sub>4</sub> phase and amorphous C can be found in the products. SEM observation shows a special micro-morphol. of sample, which is favorable for enhancement of electrochem. properties. The discharge capacity of the LiFePO<sub>4</sub>/C composite was 135 mAh/g, close to the charge capacity of 153 mAh/g at low rate of 0.1 C. At 0.2 C, the specific capacity was .apprx.117.4 mAh/g, which is satisfied for power source of Elec. Vehicle for its flat discharge platform.  
ST electrochemi iron lithium phosphate cathode composite pyrolysis carbon synthesis  
IT Battery cathodes  
    Cathodes  
    Cyclic voltammetry  
    Surface structure  
    X-ray diffraction  
        (improvement of electrochem. performance of LiFePO<sub>4</sub> cathode composite material using a in situ pyrolysis carbon synthesis procedure)  
IT 21324-40-3, Lithium hexafluorophosphate  
RL: NUU (Other use, unclassified); USES (Uses)

(electrolyte; improvement of electrochem. performance of LiFePO<sub>4</sub> cathode composite material using a in situ pyrolysis carbon synthesis procedure)

IT 7440-44-0P, Carbon, uses 15365-14-7P, Iron lithium phosphate (FeLiPO<sub>4</sub>)  
 RL: OCU (Occurrence, unclassified); PRP (Properties); SPN (Synthetic preparation); TEM (Technical or engineered material use); OCCU (Occurrence); PREP (Preparation); USES (Uses)  
 (improvement of electrochem. performance of LiFePO<sub>4</sub> cathode composite material using a in situ pyrolysis carbon synthesis procedure)

IT 516-03-0, Iron oxalate 554-13-2, Lithium carbonate 7722-76-1, Ammonium dihydrogen phosphate 9002-89-5, Polyvinyl alcohol  
 RL: RCT (Reactant); RACT (Reactant or reagent)  
 (improvement of electrochem. performance of LiFePO<sub>4</sub> cathode composite material using a in situ pyrolysis carbon synthesis procedure)

IT 107-21-1, Ethylene glycol, uses 616-38-6, Dimethyl carbonate  
 RL: NUU (Other use, unclassified); USES (Uses)  
 (solvent containing; improvement of electrochem. performance of LiFePO<sub>4</sub> cathode composite material using a in situ pyrolysis carbon synthesis procedure)

RE.CNT 7 THERE ARE 7 CITED REFERENCES AVAILABLE FOR THIS RECORD

RE

- (1) Chen, Z; J Electrochem Soc 2002, V149, PA1184 CAPLUS
- (2) Huang, H; Electrochim Solid-State Lett 2001, V4, PA170 CAPLUS
- (3) Lu, J; Acta Phys-Chim Sin 2001, V21(3), P319
- (4) Lu, J; Rare Metal Mat Eng 2001, V33(7), P679
- (5) Padhi, A; J Electrochem Soc 1997, V144, P1188 CAPLUS
- (6) Prosini, P; Electrochim Acta 2001, V46, P3517 CAPLUS
- (7) Yamada, A; J Electrochem Soc 2001, V148, PA224 CAPLUS

L81 ANSWER 9 OF 21 CAPLUS COPYRIGHT 2008 ACS on STN

AN 2007:286942 CAPLUS

DN 146:299360

ED Entered STN: 16 Mar 2007

TI Cathode material for manufacturing a rechargeable battery

IN Yang, Chih-Wei

PA Aquire Energy Co., Ltd., Taiwan

SO U.S. Pat. Appl. Publ., 17pp., Cont.-in-part of U.S. Ser. No. 222,569.  
 CODEN: USXXCO

DT Patent

LA English

INCL 429209000; 423306000; 429217000; 252182100; 429232000

CC 52-2 (Electrochemical, Radiational, and Thermal Energy Technology)  
 Section cross-reference(s): 49

FAN.CNT 8

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	US 20070059598	A1	20070315	US 2006-510096	20060825
	US 20060257307	A1	20061116	US 2005-222569	20050909
	AT 385999	T	20080315	AT 2005-256174	20051003
	US 20070207385	A1	20070906	US 2007-747746	20070511
	US 20070238021	A1	20071011	US 2007-764686	20070618
	US 20080107967	A1	20080508	US 2007-940283	20071114
	US 20080138710	A1	20080612	US 2007-940276	20071114
PRAI	US 2005-222569	A2	20050909		
	TW 2005-94115023	A	20050510		
	EP 2005-256174	A	20051003		
	CN 2006-10080365	A	20060511		
	US 2006-510096	A2	20060825		
	US 2006-518805	A2	20060911		

US 2007-747746	A2	20070511
US 2007-764629	A2	20070618
CLASS		
PATENT NO.	CLASS	PATENT FAMILY CLASSIFICATION CODES
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US 20070059598	INCL	429209000; 423306000; 429217000; 252182100; 429232000
	IPCI	C01B0025-26 [I,A]; C01B0025-00 [I,C*]; H01M0004-02 [I,A]; H01M0004-62 [I,A]
	IPCR	C01B0025-00 [I,C]; C01B0025-26 [I,A]; H01M0004-02 [I,C]; H01M0004-02 [I,A]; H01M0004-62 [I,C]; H01M0004-62 [I,A]
	NCL	429/209.000; 252/182.100; 423/306.000; 429/217.000; 429/232.000
	ECLA	H01M004/58D; H01M004/136; T01M; T01M; T01M
US 20060257307	IPCI	C01B0025-26 [I,A]; C01B0025-00 [I,C*]
	IPCR	C01B0025-00 [I,C]; C01B0025-26 [I,A]
	NCL	423/306.000
	ECLA	H01M004/58D; C01B025/45; C01D015/02; H01M004/131; H01M004/136; H01M004/485; H01M004/62; H01M004/62B; H01M004/62C2; M01P; T01M; T01M; T01M; T01M; Y01N
AT 385999	IPCI	C01G0049-00 [I,C]; C01G0049-00 [I,A]; C01B0013-14 [I,C]; C01B0013-14 [I,A]; C01B0025-00 [I,C]; C01B0025-45 [I,A]; C01D0001-00 [I,C]; C01D0001-02 [I,A]; H01M0004-48 [I,C]; H01M0004-48 [I,A]; H01M0004-58 [I,C]; H01M0004-58 [I,A]
	IPCR	C01G0049-00 [I,C]; C01G0049-00 [I,A]; C01B0013-14 [I,C]; C01B0013-14 [I,A]; C01B0025-00 [I,C]; C01B0025-45 [I,A]; C01D0001-00 [I,C]; C01D0001-02 [I,A]; H01M0004-48 [I,C]; H01M0004-48 [I,A]; H01M0004-58 [I,C]; H01M0004-58 [I,A]
	ECLA	C01B025/37; H01M004/58D; M01P; M01P; M01P; T01M; T01M
US 20070207385	IPCI	H01M0004-58 [I,A]; H01M0004-62 [I,A]
	IPCR	H01M0004-58 [I,C]; H01M0004-58 [I,A]; H01M0004-62 [I,C]; H01M0004-62 [I,A]
	NCL	429/231.900; 252/182.100; 429/231.600; 429/231.950; 429/232.000
	ECLA	H01M004/58D; H01M004/131; H01M004/136; H01M004/485; H01M004/62; H01M004/62B; H01M004/62C2; T01M; T01M; T01M; T01M; T01M
US 20070238021	IPCI	H01M0004-38 [I,A]
	IPCR	H01M0004-38 [I,C]; H01M0004-38 [I,A]
	NCL	429/218.100
	ECLA	H01M004/58D; H01M004/485; H01M004/58; H01M004/62C2; T01M; T01M; T01M
US 20080107967	IPCI	H01M0004-38 [I,A]; H01M0004-48 [I,A]
	NCL	429/219.000; 429/229.000; 429/231.600; 429/218.100; 429/231.800; 429/231.900; 429/231.950; 429/231.500; 429/223.000; 429/222.000
US 20080138710	IPCI	H01M0004-36 [I,A]; H01M0004-42 [I,A]; H01M0004-44 [I,A]; H01M0004-46 [I,A]
	NCL	429/222.000; 429/229.000; 429/231.000; 429/231.600; 252/182.330

AB A cathode material having olivine or NASICON structures and includes micrometer-sized secondary particles having a particle size of 1-50  $\mu\text{m}$ . Each of the micrometer-sized secondary particles is composed of crystalline nanometer-sized primary particles of a metal compound having a particle size of 10-500 nm. The metal compound has a formula  $A_3xM_2y(PO_4)_3$  with A being a Group IA, IIA, or IIIA element, M being a 2nd metal element from Groups IIA, IIIA, or a transition element, and  $0 < x \leq 1.2$ , and  $0 < y \leq 1.6$ . Carbon particles adhere to the surface of the crystalline nanometer-sized primary particles. The cathode material has a

BET sp. surface area of 5-100 m<sup>2</sup>/g. The cathode material is coated on an electrode plate. The cathode material contains a binder, such as styrene-butadiene rubber or polyvinylidene fluoride. The cathode material contains a thickener, especially CM-cellulose.

ST cathode material rechargeable battery lithium ferrous phosphate

IT Styrene-butadiene rubber, uses  
RL: MOA (Modifier or additive use); USES (Uses)  
(cathode containing; cathode material for manufacturing rechargeable battery)

IT Battery cathodes  
NASICONS  
Secondary batteries  
(cathode material for manufacturing rechargeable battery)

IT Fluoropolymers, uses  
RL: MOA (Modifier or additive use); TEM (Technical or engineered material use); USES (Uses)  
(cathode material for manufacturing rechargeable battery)

IT Charcoal  
RL: PEP (Physical, engineering or chemical process); PROC (Process)  
(cathode material for manufacturing rechargeable battery)

IT 7440-44-0, Carbon, uses  
RL: MOA (Modifier or additive use); TEM (Technical or engineered material use); USES (Uses)  
(anode, cathode containing; cathode material for manufacturing rechargeable battery)

IT 9004-32-4, Carboxymethyl cellulose  
RL: MOA (Modifier or additive use); USES (Uses)  
(cathode containing; cathode material for manufacturing rechargeable battery)

IT 24937-79-9, Polyvinylidene fluoride  
RL: MOA (Modifier or additive use); TEM (Technical or engineered material use); USES (Uses)  
(cathode containing; cathode material for manufacturing rechargeable battery)

IT 50-99-7, Glucose, processes 57-50-1, Sucrose, processes 77-92-9  
, Citric acid, processes 144-62-7, Oxalic acid, processes  
RL: PEP (Physical, engineering or chemical process); PROC (Process)  
(cathode material for manufacturing rechargeable battery)

IT 1310-65-2, Lithium hydroxide 7439-89-6, Iron, reactions 7446-70-0,  
Aluminum chloride, reactions 7664-38-2, Phosphoric acid, reactions  
7705-08-0, Ferric chloride, reactions 7786-30-3, Magnesium chloride,  
reactions 10421-48-4, Ferric nitrate  
RL: PEP (Physical, engineering or chemical process); RCT (Reactant); PROC  
(Process); RACT (Reactant or reagent)  
(cathode material for manufacturing rechargeable battery)

IT 15365-14-7P, Iron lithium phosphate felipo4 928163-03-5P  
RL: IMF (Industrial manufacture); PEP (Physical, engineering or chemical process); PRP (Properties); TEM (Technical or engineered material use);  
PREP (Preparation); PROC (Process); USES (Uses)  
(cathode material; cathode material for manufacturing rechargeable battery)

IT 7429-90-5, Aluminum, uses  
RL: TEM (Technical or engineered material use); USES (Uses)  
(electrode plate; cathode material for manufacturing rechargeable battery)

IT 96-49-1, Ethylene carbonate 105-58-8, Diethyl carbonate  
RL: TEM (Technical or engineered material use); USES (Uses)  
(electrolyte containing; cathode material for manufacturing rechargeable battery)

IT 21324-40-3, Lithium hexafluorophosphate  
RL: TEM (Technical or engineered material use); USES (Uses)

(electrolyte; cathode material for manufacturing rechargeable battery)

IT 9003-55-8  
 RL: MOA (Modifier or additive use); USES (Uses)  
 (styrene-butadiene rubber, cathode containing; cathode material for manufacturing rechargeable battery)

L81 ANSWER 10 OF 21 CAPLUS COPYRIGHT 2008 ACS on STN  
 AN 2007:186394 CAPLUS  
 DN 146:255322  
 ED Entered STN: 20 Feb 2007  
 TI Method for microwave synthesis of carbon coated lithium iron phosphate used as composite cathode material of lithium batteries  
 IN Zou, Zhigang; Zhu, Mei; Chen, Hong  
 PA Nanjing University, Peop. Rep. China  
 SO Faming Zhuanli Shenqing Gongkai Shuomingshu, 9pp.  
 CODEN: CNXXEV

DT Patent  
 LA Chinese  
 CC 52-2 (Electrochemical, Radiational, and Thermal Energy Technology)  
 FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI CN 1911792	A	20070214	CN 2006-10041396	20060822
PRAI CN 2006-10041396		20060822		

CLASS

PATENT NO.	CLASS	PATENT FAMILY CLASSIFICATION CODES
CN 1911792	IPCI	C01B0025-45 [I,A]; C01B0025-00 [I,C*]; H01M0004-58 [I,A] IPCR C01B0025-00 [I,C]; C01B0025-45 [I,A]

AB Carbon coated lithium iron phosphate is produced by ball milling a Li salt, a ferrous salt, a phosphate, and organic carbon source at a molar ratio of Li:Fe:P:C = 1:1:1:(0.2-2) in ethanol or acetone as a dispersant for 3-6 h, drying, tabletting, placing into alumina pot containing activated carbon, and heating under microwave irradiation for 5-12 min. The Li salt can be lithium carbonate, lithium hydroxide, lithium acetate, lithium lactate, lithium oxalate, lithium citrate, or lithium formate. The ferrous salt can be ferrous oxalate, ferrous acetate, or ferrous lactate. The phosphate can be diammonium hydrogen phosphate or ammonium dihydrogen phosphate. The organic carbon source can be citric acid, oxalic acid, tartaric acid, glucose, sucrose, lactose, or maltose. The composite material can be used to fabricate cathodes for lithium batteries.

ST carbon coated lithium iron phosphate composite cathode battery  
 IT Secondary batteries  
 (lithium; production of carbon coated lithium iron phosphate used as composite cathode material of lithium batteries)

IT Ball milling  
 Battery cathodes  
 Microwave heating  
 (production of carbon coated lithium iron phosphate used as composite cathode material of lithium batteries)

IT 64-17-5, Ethanol, uses 67-64-1, Acetone, uses  
 RL: NUU (Other use, unclassified); USES (Uses)  
 (dispersant; production of carbon coated lithium iron phosphate used as composite cathode material of lithium batteries)

IT 7440-44-0, Carbon, uses  
 RL: PEP (Physical, engineering or chemical process); TEM (Technical or engineered material use); PROC (Process); USES (Uses)  
 (lithium iron phosphate coated with, cathode material; production of carbon coated lithium iron phosphate used as composite

cathode material of lithium batteries)  
 IT 15365-14-7P, Iron lithium phosphate FeLiPO<sub>4</sub>  
 RL: IMF (Industrial manufacture); PEP (Physical, engineering or chemical process); TEM (Technical or engineered material use); PREP (Preparation); PROC (Process); USES (Uses)  
     (product of carbon coated lithium iron phosphate used as composite cathode material of lithium batteries)  
 IT 50-99-7, Glucose, processes 57-50-1, Sucrose, processes 63-42-3,  
     Lactose 69-79-4, Maltose 77-92-9, Citric acid, processes  
     87-69-4, Tartaric acid, processes 144-62-7, Oxalic acid,  
     processes  
     RL: PEP (Physical, engineering or chemical process); PROC (Process)  
     (product of carbon coated lithium iron phosphate used as composite cathode material of lithium batteries)  
 IT 516-03-0, Ferrous oxalate 546-89-4, Lithium acetate 553-91-3, Lithium oxalate 554-13-2, Lithium carbonate 556-63-8, Lithium formate 867-55-0, Lithium lactate 919-16-4, Lithium citrate 1310-65-2, Lithium hydroxide 3094-87-9, Ferrous acetate 5905-52-2, Ferrous lactate 7722-76-1, Ammonium dihydrogen phosphate 7783-28-0, Diammonium hydrogen phosphate  
     RL: PEP (Physical, engineering or chemical process); RCT (Reactant); PROC (Process); RACT (Reactant or reagent)  
     (product of carbon coated lithium iron phosphate used as composite cathode material of lithium batteries)

L81 ANSWER 11 OF 21 CAPLUS COPYRIGHT 2008 ACS on STN  
 AN 2007:152960 CAPLUS  
 DN 146:255292  
 ED Entered STN: 12 Feb 2007  
 TI Preparation of high density ultrafine composite lithium iron phosphate as cathode material for lithium ion batteries  
 IN Li, Jun; Lai, Guitang; Huang, Huimin; Xia, Xinde; Xue, Jianjun; Li, Daguang  
 PA Guangzhou Great Power Battery Co., Ltd., Peop. Rep. China  
 SO Faming Zhuanli Shenqing Gongkai Shuomingshu, 11pp.  
     CODEN: CNXXEV  
 DT Patent  
 LA Chinese  
 CC 52-2 (Electrochemical, Radiational, and Thermal Energy Technology)  
     Section cross-reference(s): 49

FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI CN 1907844	A	20070207	CN 2006-10037041	20060811
PRAI CN 2006-10037041		20060811		

CLASS

PATENT NO.	CLASS	PATENT FAMILY CLASSIFICATION CODES
CN 1907844	IPCI	C01B0025-45 [I,A]; C01B0025-00 [I,C*]; H01M0004-58 [I,A] IPCR C01B0025-00 [I,C]; C01B0025-45 [I,A]

AB The title process comprises: (1) mixing P compound, Li salt with Fe salt by a mol. ratio of 1-1.1:1-1.1:1-1.1 under the addition of compds. of doping elements or carbon organic compds. (conductive additive); (2) adding organic acid as support; (3) allowing reaction at ≤ 100 °C to obtain nanometer precursor under regulating pH to 4.0-9.0 with LiOH and ammonia liquor; and (4) calcining at 600-800 °C for 5-300 min to give the title product. The Li salt is LiOH, Li<sub>2</sub>CO<sub>3</sub>, lithium oxalate, lithium fluoride, lithium phosphate and/or lithium acetate. The Fe salt is soluble Fe salt, ferrous acetate and/or ferrous sulfate. The P compound is NH<sub>4</sub>H<sub>2</sub>PO<sub>4</sub> and/or (NH<sub>4</sub>)<sub>2</sub>HPO<sub>4</sub>. The doping compound is magnesium acetate and/or magnesium hydroxide. The organic acid is oxalic acid, tartaric acid, acrylic

acid, citric acid, polyacrylic acid, humic acid, polyvinylpyrrolidone, 2-ethylhexoic acid, and/or succinic acid. The carbon-containing organic additive is polypropylene, polyacrylamide, glucose, sucrose, and/or starch. The title product has stable structure, good thermal stability, and good recycling properties.

ST high density ultrafine composite lithium iron phosphate anode battery

IT Secondary batteries  
(lithium, lithium ion; preparation of high d. ultrafine composite lithium iron phosphate as anode material for lithium ion batteries)

IT Nanoparticles Powders  
(nanopowders; preparation of high d. ultrafine composite lithium iron phosphate as anode material for lithium ion batteries)

IT Carbon black, uses  
Fluoropolymers, uses  
Humic acids  
RL: NUU (Other use, unclassified); TEM (Technical or engineered material use); USES (Uses)  
(preparation of high d. ultrafine composite lithium iron phosphate as anode material for lithium ion batteries)

IT 9003-07-0, Celgard2300  
RL: NUU (Other use, unclassified); TEM (Technical or engineered material use); USES (Uses)  
(film; preparation of high d. ultrafine composite lithium iron phosphate as anode material for lithium ion batteries)

IT 50-99-7, Glucose, uses 57-50-1, Sucrose, uses 77-92-9, Citric acid, uses 79-10-7, Acrylic acid, uses 87-69-4, Tartaric acid, uses 96-49-1, Ethylene carbonate 110-15-6, Succinic acid, uses 142-72-3, Magnesium acetate 144-62-7, Oxalic acid, uses 149-57-5, 2-Ethylhexoic acid 546-89-4, Lithium acetate 553-91-3, Lithium oxalate 554-13-2, Lithium carbonate 616-38-6, Dimethyl carbonate 1309-42-8, Magnesium hydroxide 1310-65-2, Lithium hydroxide 1345-25-1, Ferrous oxide, uses 3094-87-9, Ferrous acetate 7429-90-5, Aluminum, uses 7439-93-2, Lithium, uses 7440-44-0, Carbon, uses 7440-50-8, Copper, uses 7664-41-7, Ammonia, uses 7720-78-7, Ferrous sulfate 7722-76-1, Ammonium dihydrogen phosphate 7783-28-0, Diammonium hydrogen phosphate 7789-24-4, Lithium fluoride, uses 9003-01-4, Polyacrylic acid 9003-05-8, Polyacrylamide 9003-39-8, Polyvinylpyrrolidone 9005-25-8, Starch, uses 10377-52-3, Lithium phosphate 15365-14-7, Iron lithium phosphate FeLiPO<sub>4</sub> 21324-40-3, Lithium hexafluorophosphate 24937-79-9, Polyvinylidene fluoride  
RL: NUU (Other use, unclassified); TEM (Technical or engineered material use); USES (Uses)  
(preparation of high d. ultrafine composite lithium iron phosphate as anode material for lithium ion batteries)

L81 ANSWER 12 OF 21 CAPLUS COPYRIGHT 2008 ACS on STN  
AN 2006:1216545 CAPLUS  
DN 147:238626  
ED Entered STN: 20 Nov 2006  
TI Mass and charge transport in hierarchically organized storage materials.  
Example: Porous active materials with nanocoated walls of pores  
AU Gaberscek, Miran; Dominko, Robert; Bele, Marjan; Remskar, Maja; Jamnik, Janez  
CS National Institute of Chemistry, Ljubljana, SI-1000, Slovenia  
SO Solid State Ionics (2006), 177(35-36), 3015-3022  
CODEN: SSIOD3; ISSN: 0167-2738  
PB Elsevier B.V.  
DT Journal

LA English  
CC 52-2 (Electrochemical, Radiational, and Thermal Energy Technology)  
Section cross-reference(s): 72

AB To enhance the kinetics of poorly conducting cathode materials for Li batteries, the authors have proposed a number of strategies based on crushing the active material into nanopowder and embedding the powder into a carbon-based web or coating. Using the well-elaborated example of LiFePO<sub>4</sub>, we demonstrate that the same goal can be achieved with a different approach where the active material remains in a form of large (1-20  $\mu\text{m}$ ) single crystals. Instead of crushing the material, we make it porous, with average pore size around 50 nm and pore surface area of 25 m<sup>2</sup>/g. The walls of the pores (but also the outer surfaces of crystals) are covered with ca. 1 nm thick carbon film. Most surprisingly, such a unique nanoarchitecture can be prepared using a simple sol-gel based procedure including a single heat treatment. The crucial part is the selection of appropriate carbon precursor. For example, citric acid decomp. quite vigorously into gases and solid carbon at temps. up to ca. 450 °C. This range matches exactly the first solidification of LiFePO<sub>4</sub>. Thus, the evolving gases can create an interconnected web of pores while the solid parts (carbon) are deposited simultaneously on the walls of pores. We further show that a carbon content of less than 3% is already sufficient for surpassing the percolation threshold with respect to surface conductivity of carbon. Using more carbon can decrease the rate performance so a fine balance is required in this respect. Most importantly, carbonization at a temperature of slightly less than 700 °C is sufficient to achieve a composite conductivity of the order of 10<sup>-2</sup> S cm<sup>-2</sup>-more than sufficient for good cathode kinetics. In the end, we show new evidence that the phase that is responsible for high conductivity of LiFePO<sub>4</sub>-C composites is indeed the carbon phase.

ST lithium iron phosphate carbon composite lithium battery cathode;  
titania carbon composite lithium battery cathode

IT Secondary batteries  
(lithium; mass and charge transport in porous active materials with nanocoated walls of pores, hierarchically organized storage materials)

IT Composites  
Porous materials  
(mass and charge transport in porous active materials with nanocoated walls of pores, hierarchically organized storage materials)

IT 13463-67-7, Titania, reactions  
RL: PEP (Physical, engineering or chemical process); RCT (Reactant); PROC (Process); RACT (Reactant or reagent)  
(composite with C; mass and charge transport in porous active materials with nanocoated walls of pores, hierarchically organized storage materials)

IT 15365-14-7, Iron lithium phosphate felipo4  
RL: TEM (Technical or engineered material use); USES (Uses)  
(composite with C; mass and charge transport in porous active materials with nanocoated walls of pores, hierarchically organized storage materials)

IT 7440-44-0, Carbon, uses  
RL: TEM (Technical or engineered material use); USES (Uses)  
(composite with FeLiPO<sub>4</sub>; mass and charge transport in porous active materials with nanocoated walls of pores, hierarchically organized storage materials)

IT 77-92-9, Citric acid, reactions 3522-50-7, Ferric citrate  
7664-38-2, Phosphoric acid, reactions 10377-52-3, Lithium phosphate  
13453-80-0, Lithium dihydrogen phosphate  
RL: PEP (Physical, engineering or chemical process); RCT (Reactant); PROC (Process); RACT (Reactant or reagent)  
(mass and charge transport in porous active materials with nanocoated walls of pores, hierarchically organized storage materials)

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- (2) Chen, Z; J Electrochim Soc 2002, V149, PA1184 CAPLUS
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L81 ANSWER 13 OF 21 CAPLUS COPYRIGHT 2008 ACS on STN

AN 2006:958597 CAPLUS

DN 146:503916

ED Entered STN: 18 Sep 2006

TI Synthesis and effect of forming Fe2P phase on the physics and electrochemical properties of LiFePO4/C materials

AU Xu, Yanbin; Lu, Yingjun; Yan, Lan; Yang, Zhengyin; Yang, Rudong

CS College of Chemistry and Chemical Engineering, Lanzhou University, Lanzhou, 730000, Peop. Rep. China

SO Journal of Power Sources (2006), 160(1), 570-576

CODEN: JPSODZ; ISSN: 0378-7753

PB Elsevier B.V.

DT Journal

LA English

CC 52-2 (Electrochemical, Radiational, and Thermal Energy Technology)  
Section cross-reference(s): 72

AB A series of LiFePO4/C materials were prepared by a reformative solid-coordination method which uses citric acid as the coordination agent and C source. A monodentate coordination bond of -COO-M forms gradually and that helps to disperse Li<sup>+</sup> or Fe<sup>2+</sup> in the homogeneous gel during grinding. Impure phase Fe2P was detected in the LiFePO4/C composites with increasing annealing temperature. The remnant coating C was considered to be

the  
reductive in pure N. The amts. of C, particle size and morphol. were studied in detail and all the results can be related to the formation of Fe2P. The electro-conductive Fe2P phase in LiFePO4/C composites has an important role in increasing electronic conductivity and it improves the electrochem. performance of LiFePO4/C including the polarization phenomenon, comparatively high reversible capability, stable cycling performance and slight trend of less loss of rate capability.

ST iron phosphide lithium iron phosphate carbon cathode lithium battery

IT Secondary batteries

(lithium; synthesis of LiFePO4/C cathode material for lithium batteries with forming of Fe2P phase and its effects)

IT Battery cathodes

(synthesis of LiFePO4/C cathode material for lithium batteries with forming of Fe2P phase and its effects)

IT 77-92-9, Citric acid, processes

RL: PEP (Physical, engineering or chemical process); PROC (Process)

(in synthesis of LiFePO<sub>4</sub>/C cathode material for lithium batteries with forming of Fe<sub>2</sub>P phase and its effects)

IT 1310-43-6, Iron phosphide (Fe<sub>2</sub>P)  
RL: FMU (Formation, unclassified); FORM (Formation, nonpreparative) (synthesis of LiFePO<sub>4</sub>/C cathode material for lithium batteries with forming of Fe<sub>2</sub>P phase and its effects)

IT 7440-44-0P, Carbon, uses 15365-14-7P, Iron lithium phosphate (FeLiPO<sub>4</sub>)  
RL: SPN (Synthetic preparation); TEM (Technical or engineered material use); PREP (Preparation); USES (Uses) (synthesis of LiFePO<sub>4</sub>/C cathode material for lithium batteries with forming of Fe<sub>2</sub>P phase and its effects)

RE.CNT 24 THERE ARE 24 CITED REFERENCES AVAILABLE FOR THIS RECORD

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(24) Yang, S; J Power Sources 2003, V119-121, P239 CAPLUS

L81 ANSWER 14 OF 21 CAPLUS COPYRIGHT 2008 ACS on STN  
AN 2005:1339204 CAPLUS  
DN 145:359878  
ED Entered STN: 26 Dec 2005  
TI Synthesis, characterization and properties of LiFePO<sub>4</sub>/C cathode material  
AU Zhou, Xin-wen; Zhan, Dan; Wang, Li-na; Liu, Qiao-yun; Zong, Hong-xing; Zhang, Ke-li  
CS College of Chemistry and Molecular Sciences, Wuhan University, Wuhan, 430072, Peop. Rep. China  
SO Wuhan University Journal of Natural Sciences (2005), 10(5), 909-912  
CODEN: WUNSFW; ISSN: 1007-1202  
PB Wuhan University Journals Press  
DT Journal  
LA English  
CC 52-2 (Electrochemical, Radiational, and Thermal Energy Technology)  
AB Lithium iron phosphate coated with carbon (LiFePO<sub>4</sub>/C) was synthesized by a rheol. phase reaction using a comparatively lower temperature and less sintering time. The carbon came from citric acid, which acted as a new carbon

source. It was characterized by thermogravimetry and DTA (TG/DTA), X ray diffractometer (XRD), Element Anal. (EA) and Scanning electron microscope (SEM). We also studied the electrochem. properties of the material. The first discharge capacity of the LiFePO<sub>4</sub>/C is 121 mAh · g<sup>-1</sup> at 10 mA · g<sup>-1</sup> at room temperature When the c.d. increased to 100 mA · g<sup>-1</sup> the first discharge capacity decreased to 110 mAh · g<sup>-1</sup> and retained 95% of the initial capacity after 100 cycles. The LiFePO<sub>4</sub>/C obtained shows a good electrochem. capacity and cycle ability at a large c.d.

ST lithium iron phosphate lithium ion battery cathode  
IT Secondary batteries  
    (lithium, lithium ion; synthesis, characterization and properties of LiFePO<sub>4</sub>/C cathode material for lithium ion batteries)  
IT Battery cathodes  
Electrochemistry  
Surface structure  
    (synthesis, characterization and properties of LiFePO<sub>4</sub>/C cathode material for lithium ion batteries)  
IT 77-92-9, Citric acid, reactions  
RL: RCT (Reactant); RACT (Reactant or reagent)  
    (carbon source; synthesis, characterization and properties of LiFePO<sub>4</sub>/C cathode material for lithium ion batteries)  
IT 7440-44-0P, Carbon, uses  
RL: DEV (Device component use); SPN (Synthetic preparation); PREP (Preparation); USES (Uses)  
    (coated onto LiFePO<sub>4</sub>; synthesis, characterization and properties of LiFePO<sub>4</sub>/C cathode material for lithium ion batteries)  
IT 15365-14-7P, Iron lithium phosphate felipo<sub>4</sub>  
RL: DEV (Device component use); SPN (Synthetic preparation); PREP (Preparation); USES (Uses)  
    (coated with carbon; synthesis, characterization and properties of LiFePO<sub>4</sub>/C cathode material for lithium ion batteries)

RE.CNT 16 THERE ARE 16 CITED REFERENCES AVAILABLE FOR THIS RECORD

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L81 ANSWER 15 OF 21 CAPLUS COPYRIGHT 2008 ACS on STN

AN 2005:673860 CAPLUS

DN 143:176223

ED Entered STN: 31 Jul 2005

TI Composite mixed oxides as active battery electrodes, especially for rechargeable lithium batteries

IN Gauthier, Gilles; Le Cras, Frederic; Lignier, Helene; Gabelle, Jean Louis  
PA Commissariat a l'Energie Atomique, Fr.

SO Fr. Demande, 45 pp.

CODEN: FRXXBL

DT Patent  
 LA French  
 IC ICM H01M004-60  
 ICS H01M004-26  
 CC 52-2 (Electrochemical, Radiational, and Thermal Energy Technology)  
 FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	FR 2865576	A1	20050729	FR 2004-50156	20040128
	FR 2865576	B1	20060428		
	WO 2005076390	A2	20050818	WO 2005-FR50045	20050126
	WO 2005076390	A3	20051006		
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW RW: BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG					
EP 1709702	A2	20061011	EP 2005-717687	20050126	
EP 1709702	B1	20071205			
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, FI, RO, CY, TR, BG, CZ, EE, HU, PL, SK, IS					
JP 2007520038	T	20070719	JP 2006-550260	20050126	
AT 380399	T	20071215	AT 2005-717687	20050126	
ES 2299019	T3	20080516	ES 2005-717687	20050126	
US 20070152185	A1	20070705	US 2007-586601	20070104	
PRAI FR 2004-50156	A	20040128			
WO 2005-FR50045	W	20050126			

#### CLASS

	PATENT NO.	CLASS	PATENT FAMILY CLASSIFICATION CODES	
WO 2005076390	FR 2865576	ICM	H01M004-60	
		ICS	H01M004-26	
		IPCI	H01M0004-36 [I,C]; H01M0004-26 [I,C]; H01M0004-60 [I,A]; H01M0004-26 [I,A]	
		IPCR	H01M0004-36 [I,C]; H01M0004-60 [I,A]; C01G0001-02 [I,C*]; C01G0001-02 [I,A]; C01G0001-06 [I,C*]; C01G0001-06 [I,A]; H01M0004-26 [I,C]; H01M0004-26 [I,A]; H01M0004-48 [I,C*]; H01M0004-48 [I,A]; H01M0004-50 [N,C*]; H01M0004-50 [N,A]; H01M0004-52 [N,C*]; H01M0004-52 [N,A]; H01M0004-58 [I,C*]; H01M0004-58 [I,A]; H01M0004-62 [I,C*]; H01M0004-62 [I,A]	
		IPCR	C01G0001-02 [I,C*]; C01G0001-02 [I,A]; C01G0001-06 [I,C*]; C01G0001-06 [I,A]; H01M0004-48 [I,C*]; H01M0004-48 [I,A]; H01M0004-50 [N,C*]; H01M0004-50 [N,A]; H01M0004-52 [N,C*]; H01M0004-52 [N,A]; H01M0004-58 [I,C*]; H01M0004-58 [I,A]; H01M0004-62 [I,C*]; H01M0004-62 [I,A]	
		ECLA	C01G001/02; C01G001/06; H01M004/48B2; H01M004/62C2; M01P; T01M; T01M; T01M	
		IPCI	H01M0004-04 [I,C]; H01M0004-04 [I,A]; H01M0004-58 [I,C]; H01M0004-58 [I,A]	
		IPCR	H01M0004-04 [I,C]; H01M0004-04 [I,A]; C01G0001-02 [I,C*]; C01G0001-02 [I,A]; C01G0001-06 [I,C*]; C01G0001-06 [I,A]; H01M0004-48 [I,C*]; H01M0004-48 [I,A]; H01M0004-50 [N,C*]; H01M0004-50 [N,A];	
	EP 1709702			

		H01M0004-52 [N,C*]; H01M0004-52 [N,A]; H01M0004-58 [I,C]; H01M0004-58 [I,A]; H01M0004-62 [I,C*]; H01M0004-62 [I,A]
	ECLA	C01G001/02; C01G001/06; H01M004/48B2; H01M004/62C2; M01P; T01M; T01M; T01M
JP 2007520038	IPCI	H01M0004-04 [I,A]; H01M0004-58 [I,A]; H01M0004-62 [I,A]; H01M0010-40 [N,A]; H01M0010-36 [N,C*]
	IPCR	H01M0004-04 [I,C]; H01M0004-04 [I,A]; C01G0001-02 [I,C*]; C01G0001-02 [I,A]; C01G0001-06 [I,C*]; C01G0001-06 [I,A]; H01M0004-48 [I,C*]; H01M0004-48 [I,A]; H01M0004-50 [N,C*]; H01M0004-50 [N,A]; H01M0004-52 [N,C*]; H01M0004-52 [N,A]; H01M0004-58 [I,C]; H01M0004-58 [I,A]; H01M0004-62 [I,C]; H01M0004-62 [I,A]; H01M0010-36 [N,C]; H01M0010-40 [N,A]
	ECLA	C01G001/02; C01G001/06; H01M004/48B2; H01M004/62C2; M01P; T01M; T01M
	FTERM	5H029/AJ03; 5H029/AJ12; 5H029/AJ14; 5H029/AK03; 5H029/AL06; 5H029/AL07; 5H029/AL08; 5H029/CJ02; 5H029/CJ08; 5H029/CJ28; 5H029/DJ08; 5H029/EJ04; 5H029/EJ11; 5H029/EJ12; 5H029/HJ01; 5H029/HJ02; 5H029/HJ05; 5H029/HJ07; 5H029/HJ14; 5H050/AA08; 5H050/AA15; 5H050/AA19; 5H050/BA16; 5H050/BA17; 5H050/CA07; 5H050/CA08; 5H050/CA09; 5H050/CB07; 5H050/CB08; 5H050/CB09; 5H050/DA09; 5H050/EA08; 5H050/EA22; 5H050/EA23; 5H050/GA02; 5H050/GA10; 5H050/GA27; 5H050/HA01; 5H050/HA02; 5H050/HA05; 5H050/HA07; 5H050/HA14; 5H050/HA20
AT 380399	IPCI	H01M0004-04 [I,C]; H01M0004-04 [I,A]; H01M0004-58 [I,C]; H01M0004-58 [I,A]
	IPCR	H01M0004-04 [I,C]; H01M0004-04 [I,A]; C01G0001-02 [I,C*]; C01G0001-02 [I,A]; C01G0001-06 [I,C*]; C01G0001-06 [I,A]; H01M0004-48 [I,C*]; H01M0004-48 [I,A]; H01M0004-50 [N,C*]; H01M0004-50 [N,A]; H01M0004-52 [N,C*]; H01M0004-52 [N,A]; H01M0004-58 [I,C]; H01M0004-58 [I,A]; H01M0004-62 [I,C*]; H01M0004-62 [I,A]
	ECLA	C01G001/02; C01G001/06; H01M004/48B2; H01M004/62C2; M01P; T01M; T01M
ES 2299019	IPCI	H01M0004-04 [I,C]; H01M0004-04 [I,A]; H01M0004-58 [I,C]; H01M0004-58 [I,A]
	IPCR	H01M0004-04 [I,C]; H01M0004-04 [I,A]; C01G0001-02 [I,C*]; C01G0001-02 [I,A]; C01G0001-06 [I,C*]; C01G0001-06 [I,A]; H01M0004-48 [I,C*]; H01M0004-48 [I,A]; H01M0004-50 [N,C*]; H01M0004-50 [N,A]; H01M0004-52 [N,C*]; H01M0004-52 [N,A]; H01M0004-58 [I,C]; H01M0004-58 [I,A]; H01M0004-62 [I,C*]; H01M0004-62 [I,A]
	ECLA	C01G001/02; C01G001/06; H01M004/48B2; H01M004/62C2; M01P; T01M; T01M
US 20070152185	IPCI	H01B0001-06 [I,A]; H01B0001-18 [I,A]; H01B0001-14 [I,C*]
	NCL	252/182.100; 252/506.000; 252/507.000

AB Electrode-active materials, especially for alkali ion insertion (i.e., Na<sup>+</sup> and Li<sup>+</sup>) for lithium batteries, contain, as an active component, a composition of general formula AaDdMmZsOoNnFf,, in which: (1) A is an alkali metal, (2) D is an alkaline earth metal or a Group IIIA element, with the exception of B, (3) M is a transition metal, (4) Z is a non-metal selected from S, Se, P, As, Si, Ge, Sn, and B, (5) O is oxygen, N is nitrogen, and F is fluorine, and (6) a, d, m, z, o, n, and f are ≥0. The compns., which also contain an electron conductor, such as carbon, are prepared by thermal decomposition of homogeneously mixed precursors, which are organic or organometallic derivs. (preferably at 200–600°). Preferred

components include: (1) A = Li, Na, and K, (2) D is Mg, Al, and Ga, (3) M = Fe, Ni, Co, Mn, V, Mo, Nb, W, and Ti; preferred components are LiFePO<sub>4</sub>, LiFeBO<sub>3</sub>, or NaFeBO<sub>3</sub>.

ST electrode mixed oxide lithium rechargeable battery; iron lithium borate secondary battery electrode

IT Transition metal oxides  
RL: DEV (Device component use); TEM (Technical or engineered material use); USES (Uses)  
(alkaline earth oxides, electrode active materials; composite mixed oxides as active battery electrodes, especially for rechargeable lithium batteries)

IT Transition metal oxides  
RL: DEV (Device component use); TEM (Technical or engineered material use); USES (Uses)  
(alkali metal oxides, electrode active materials; composite mixed oxides as active battery electrodes, especially for rechargeable lithium batteries)

IT Battery electrodes  
(composite mixed oxides as active battery electrodes, especially for rechargeable lithium batteries)

IT Carboxylic acids, processes  
RL: CPS (Chemical process); PEP (Physical, engineering or chemical process); PROC (Process)  
(dicarboxylic, metal salts and complexes, electrode precursors; thermal decomposition of; composite mixed oxides as active battery electrodes, especially for rechargeable lithium batteries)

IT Carboxylic acids, processes  
RL: CPS (Chemical process); PEP (Physical, engineering or chemical process); PROC (Process)  
(hydroxy, metal salts and complexes, electrode precursors; thermal decomposition of; composite mixed oxides as active battery electrodes, especially for rechargeable lithium batteries)

IT Amino acids, processes  
Polyoxyalkylenes, processes  
RL: CPS (Chemical process); PEP (Physical, engineering or chemical process); PROC (Process)  
(metal salts and complexes, electrode precursors; thermal decomposition of; composite mixed oxides as active battery electrodes, especially for rechargeable lithium batteries)

IT Carboxylic acids, processes  
RL: CPS (Chemical process); PEP (Physical, engineering or chemical process); PROC (Process)  
(oxo, metal salts and complexes, electrode precursors; thermal decomposition of; composite mixed oxides as active battery electrodes, especially for rechargeable lithium batteries)

IT Alkali metal oxides  
Alkaline earth oxides  
Group IIIA element oxides  
RL: DEV (Device component use); TEM (Technical or engineered material use); USES (Uses)  
(transition metal oxides, electrode active materials; composite mixed oxides as active battery electrodes, especially for rechargeable lithium batteries)

IT 7440-44-0, Carbon, uses  
RL: TEM (Technical or engineered material use); USES (Uses)  
(elec. conductor; composite mixed oxides as active battery electrodes, especially for rechargeable lithium batteries)

IT 15365-14-7, Iron lithium phosphate (FeLiPO<sub>4</sub>) 332079-85-3, Iron lithium borate (FeLiBO<sub>3</sub>) 861001-97-0  
RL: DEV (Device component use); TEM (Technical or engineered material use); USES (Uses)  
(electrode active materials; composite mixed oxides as active

IT battery electrodes, especially for rechargeable lithium batteries)  
50-21-5D, Lactic acid, metal salts and complexes 56-40-6D  
, Aminoacetic acid, metal salts and complexes 56-41-7D, Alanine,  
metal salts and complexes 56-84-8D, L-Aspartic acid, metal salts  
and complexes 56-86-0D, L-Glutamic acid, metal salts and  
complexes 56-87-1D, L-Lysine, metal salts and complexes  
61-90-5D, L-Leucine, metal salts and complexes 70-26-8D,  
Ornithine, metal salts and complexes 74-79-3D, L-Arginine, metal  
salts and complexes 77-92-9D, Citric acid, metal salts and  
complexes 79-14-1D, Glycolic acid, metal salts and complexes  
87-69-4D, Tartaric acid, metal salts and complexes  
90-64-2D, Mandelic acid, metal salts and complexes  
107-21-1D, Ethylene glycol, metal salts and complexes  
110-15-6D, Succinic acid, metal salts and complexes  
110-16-7D, Maleic acid, metal salts and complexes  
110-17-8D, Fumaric acid, metal salts and complexes  
110-94-1D, Glutaric acid, metal salts and complexes  
111-46-6D, Diethylene glycol, metal salts and complexes  
123-76-2D, Levulinic acid, metal salts and complexes  
124-04-9D, Adipic acid, metal salts and complexes  
127-17-3D, Pyruvic acid, metal salts and complexes  
141-82-2D, Malonic acid, metal salts and complexes  
144-62-7D, Oxalic acid, metal salts and complexes  
298-12-4D, Glyoxylic acid, metal salts and complexes  
498-23-7D, Citraconic acid, metal salts and complexes  
499-12-7D, Aconitic acid, metal salts and complexes  
6915-15-7D, Malic acid, metal salts and complexes  
28854-76-4D, metal salts and complexes 35054-79-6D,  
Hydroxybutyric acid, metal salts and complexes 111937-70-3D,  
Hydroxyacrylic acid, metal salts and complexes 151677-68-8  
RL: CPS (Chemical process); PEP (Physical, engineering or chemical  
process); PROC (Process)  
(electrode precursors; thermal decomposition of; composite mixed  
oxides as active battery electrodes, especially for rechargeable  
lithium batteries)

RE.CNT 7 THERE ARE 7 CITED REFERENCES AVAILABLE FOR THIS RECORD

RE

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L81 ANSWER 16 OF 21 CAPLUS COPYRIGHT 2008 ACS on STN  
AN 2005:667369 CAPLUS

DN 143:389692

ED Entered STN: 29 Jul 2005

TI Porous, carbon-decorated LiFePO<sub>4</sub> prepared by sol-gel method based on  
citric acid

AU Gaberscek, Miran; Dominko, Robert; Bele, Marjan; Remskar, Maja; Hanzel,  
Darko; Jamnik, Janko

CS National Institute of Chemistry, Ljubljana, SI-1001, Slovenia

SO Solid State Ionics (2005), 176(19-22), 1801-1805

CODEN: SSIOD3; ISSN: 0167-2738

PB Elsevier B.V.

DT Journal

LA English

CC 52-2 (Electrochemical, Radiational, and Thermal Energy Technology)

AB LiFePO<sub>4</sub>-C composite cathode material for lithium batteries was

prepared. The active component consists of micrometer-sized particles having pores with a large size distribution. When filled with electrolyte, the pores are responsible for supply of ions while the distance between the pores (30-150 nm) dets. the solid-state diffusion kinetics. The walls of pores are covered with a C layer which serves as an electron conductor and is thin enough (2-3 nm) to allow penetration of Li ions. The synthesis is sol-gel based with a single heating step. The electrochem. performance is the best known for LiFePO<sub>4</sub> cathodes.

ST porous iron lithium phosphate carbon composite cathode lithium battery

IT Secondary batteries  
(lithium; porous carbon-LiFePO<sub>4</sub> composite cathode material for lithium batteries)

IT Battery cathodes  
Porous materials  
(porous carbon-LiFePO<sub>4</sub> composite cathode material for lithium batteries)

IT Sol-gel processing  
(porous carbon-LiFePO<sub>4</sub> composite cathode material prepared by sol-gel processing based on citric acid)

IT 7440-44-0, Carbon, uses  
RL: DEV (Device component use); USES (Uses)  
(porous carbon-LiFePO<sub>4</sub> composite cathode material for lithium batteries)

IT 15365-14-7P, Iron lithium phosphate (FeLiPO<sub>4</sub>)  
RL: DEV (Device component use); SPN (Synthetic preparation); PREP (Preparation); USES (Uses)  
(porous carbon-LiFePO<sub>4</sub> composite cathode material for lithium batteries)

IT 77-92-9, Citric acid, processes  
RL: CPS (Chemical process); PEP (Physical, engineering or chemical process); PROC (Process)  
(porous carbon-LiFePO<sub>4</sub> composite cathode material prepared by sol-gel processing based on citric acid)

RE.CNT 19 THERE ARE 19 CITED REFERENCES AVAILABLE FOR THIS RECORD

RE

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L81 ANSWER 17 OF 21 CAPLUS COPYRIGHT 2008 ACS on STN  
AN 2004:1028841 CAPLUS  
DN 142:222476

ED Entered STN: 01 Dec 2004  
TI Low temperature preparation of optimized phosphates for Li-battery applications  
AU Delacourt, Charles; Wurm, Calin; Reale, Priscilla; Morcrette, Mathieu; Masquelier, Christian  
CS Laboratoire de Reactivite et de Chimie des Solides, CNRS UMR 6007, Universite de Picardie Jules Verne, Amiens, 80039, Fr.  
SO Solid State Ionics (2004), 173(1-4), 113-118  
CODEN: SSIOD3; ISSN: 0167-2738  
PB Elsevier B.V.  
DT Journal  
LA English  
CC 52-2 (Electrochemical, Radiational, and Thermal Energy Technology)  
Section cross-reference(s): 49, 72  
AB The authors describe the thermodn. and kinetics that govern the precipitation of pure powders of phosphates phases of interest for Li-battery applications. The authors found precise procedures for the formation of three distinct crystalline forms of FePO<sub>4</sub>·2H<sub>2</sub>O and of pure LiMnPO<sub>4</sub>. The attempts to precipitate either LiCoPO<sub>4</sub> or LiFePO<sub>4</sub>, however, failed. In this latter case, optimized electrodes for battery applications were synthesized through a chemical conductive carbon coating at the surface of LiFePO<sub>4</sub> prepared by evaporation of an FeIII-containing aqueous solution  
ST temp optimized phosphate lithium battery electrode material  
pptn; pptn solv equil iron lithium phosphate manganese cobalt evapn;  
battery electrode lithiation phosphate capacitance potential  
carbonaceous coated  
IT Electric conductors  
(carbon; low temperature preparation of optimized phosphates for Li-battery applications)  
IT Electric potential  
(during galvanic cycling of lithium transition metal phosphates; low temperature preparation of optimized phosphates for Li-battery applications)  
IT Electric capacitance  
(galvanic cycling of lithium transition metal phosphates; low temperature preparation of optimized phosphates for Li-battery applications)  
IT Precipitation (chemical)  
(kinetics; low temperature preparation of optimized phosphates for Li-battery applications)  
IT Lithiation  
(lithium insertion; low temperature preparation of optimized phosphates for Li-battery applications)  
IT Battery electrodes  
Evaporation  
(low temperature preparation of optimized phosphates for Li-battery applications)  
IT Solubility  
(thermodn. calcns. for phases and precipitation process; low temperature preparation of optimized phosphates for Li-battery applications)  
IT 15365-14-7P, Iron lithium phosphate (FeLiPO<sub>4</sub>)  
RL: SPN (Synthetic preparation); PREP (Preparation)  
(carbonaceous material-coated; low temperature preparation of optimized phosphates for Li-battery applications)  
IT 14567-75-0P, Metastrengite  
RL: PRP (Properties); RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)  
(crystal types I and II; low temperature preparation of optimized phosphates for Li-battery applications)

IT 14986-93-7, Manganese phosphate (MnPO<sub>4</sub>)  
RL: FMU (Formation, unclassified); FORM (Formation, nonpreparative)  
(delithiated form; low temperature preparation of optimized phosphates for  
Li-battery applications)

IT 13463-10-0P, Iron phosphate (FePO<sub>4</sub>) dihydrate  
RL: CPS (Chemical process); PEP (Physical, engineering or chemical  
process); PRP (Properties); SPN (Synthetic preparation); PREP  
(Preparation); PROC (Process)  
(low temperature preparation of optimized phosphates for Li-battery  
applications)

IT 7440-44-0, Carbon, formation (nonpreparative)  
RL: FMU (Formation, unclassified); FORM (Formation, nonpreparative)  
(low temperature preparation of optimized phosphates for Li-battery  
applications)

IT 77-92-9, Citric acid, uses 107-21-1, Ethylene glycol,  
uses 1310-73-2, Sodium hydroxide, uses  
RL: MOA (Modifier or additive use); USES (Uses)  
(low temperature preparation of optimized phosphates for Li-battery  
applications)

IT 13824-49-2P, Strengite  
RL: PRP (Properties); RCT (Reactant); SPN (Synthetic preparation); PREP  
(Preparation); RACT (Reactant or reagent)  
(low temperature preparation of optimized phosphates for Li-battery  
applications)

IT 13826-59-0P, Lithium manganese phosphate (LiMnPO<sub>4</sub>)  
RL: PRP (Properties); SPN (Synthetic preparation); PREP (Preparation)  
(low temperature preparation of optimized phosphates for Li-battery  
applications)

IT 7664-38-2, Phosphoric acid, reactions 7705-08-0, Iron chloride,  
reactions 10421-48-4, Ferric nitrate 13453-80-0, Lithium dihydrogen  
phosphate  
RL: RCT (Reactant); RACT (Reactant or reagent)  
(low temperature preparation of optimized phosphates for Li-battery  
applications)

IT 36550-56-8P  
RL: SPN (Synthetic preparation); PREP (Preparation)  
(low temperature preparation of optimized phosphates for Li-battery  
applications)

IT 10377-52-3, Trilithium phosphate  
RL: FMU (Formation, unclassified); FORM (Formation, nonpreparative)  
(phase formed during some pptns.; low temperature preparation of optimized  
phosphates for Li-battery applications)

IT 12672-51-4, Cobalt hydroxide 18933-05-6, Manganese hydroxide  
51349-94-1, Manganese hydrogen phosphate  
RL: PRP (Properties)  
(solubility calcns. including; low temperature preparation of optimized  
phosphates for  
Li-battery applications)

RE.CNT 27 THERE ARE 27 CITED REFERENCES AVAILABLE FOR THIS RECORD

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L81 ANSWER 18 OF 21 CAPLUS COPYRIGHT 2008 ACS on STN  
AN 2004:689393 CAPLUS  
DN 141:352642  
ED Entered STN: 24 Aug 2004  
TI Synthesis and characterization of nano-sized LiFePO<sub>4</sub> cathode materials prepared by a citric acid-based sol-gel route  
AU Hsu, Kuei-Feng; Tsay, Sun-Yuan; Hwang, Bing-Joe  
CS Department of Chemical Engineering, National Cheng Kung University, Tainan, 701, Taiwan  
SO Journal of Materials Chemistry (2004), 14(17), 2690-2695  
CODEN: JMACEP; ISSN: 0959-9428  
PB Royal Society of Chemistry  
DT Journal  
LA English  
CC 52-2 (Electrochemical, Radiational, and Thermal Energy Technology)  
AB LiFePO<sub>4</sub>/carbon composite cathode materials were synthesized by a sol-gel process. The citric acid in the developed sol-gel process plays the role not only as a complexing agent but also as a carbon source, which improves the conductivity of the composites and hinders the growth of LiFePO<sub>4</sub> particles. Nanosized LiFePO<sub>4</sub> particles without the impurity phase were successfully synthesized. The grain size of LiFePO<sub>4</sub> particles in the range of 20-30 nm is obtained at calcining temps. 450-850°. Increasing the calcination temperature leads to a decrease in the carbon content but an increase in the conductivity of the composites in the range of 400-850°. However, the conductivity slightly decreases if the calcination temperature further increases to 950°. The LiFePO<sub>4</sub>/carbon composite synthesized at 850° shows the highest conductivity (10<sup>-3</sup> S/cm), the highest sp. capacity, and the best rate capability among the synthesized materials. It is worthy to note that the cell performance of the LiFePO<sub>4</sub> depends on the electrochem. cycling procedure employed.  
ST iron lithium phosphate carbon composite cathode synthesis sol gel; citric acid iron lithium phosphate carbon composite cathode synthesis; battery iron lithium phosphate carbon composite cathode  
IT Battery cathodes  
(synthesis and characterization of nanosized iron lithium phosphate/carbon composite cathode materials prepared by citric acid-based sol-gel method)  
IT 15365-14-7, Iron lithium phosphate (FeLiPO<sub>4</sub>)  
RL: CPS (Chemical process); DEV (Device component use); PEP (Physical, engineering or chemical process); PROC (Process); USES (Uses)  
(composite with carbon; synthesis and characterization of nanosized iron lithium phosphate/carbon composite cathode materials prepared by citric acid-based sol-gel method)  
IT 7440-44-0, Carbon, uses

RL: CPS (Chemical process); DEV (Device component use); PEP (Physical, engineering or chemical process); PROC (Process); USES (Uses)  
(composite with iron lithium phosphate; synthesis and characterization of nanosized iron lithium phosphate/carbon composite cathode materials prepared by citric acid-based sol-gel method)

IT 77-92-9, Citric acid, uses

RL: NUU (Other use, unclassified); USES (Uses)  
(synthesis and characterization of nanosized iron lithium phosphate/carbon composite cathode materials prepared by citric acid-based sol-gel method)

RE.CNT 20 THERE ARE 20 CITED REFERENCES AVAILABLE FOR THIS RECORD

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L81 ANSWER 19 OF 21 CAPLUS COPYRIGHT 2008 ACS on STN

AN 2004:3211 CAPLUS

DN 140:44760

ED Entered STN: 02 Jan 2004

TI Process for production of carbon-coated lithium-containing powders for lithium secondary battery

IN Audemer, Albane; Wurm, Calin; Morcrette, Mathieu; Gwizdala, Sylvain; Masquelier, Christian

PA Umicore, Belg.; Le Centre National de la Recherche Scientifique

SO PCT Int. Appl., 18 pp.

CODEN: PIXXD2

DT Patent

LA English

IC ICM H01M004-58

CC 52-2 (Electrochemical, Radiational, and Thermal Energy Technology)

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2004001881	A2	20031231	WO 2003-EP6628	20030619
	WO 2004001881	A3	20041229		
	W:	AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW			
	RW:	GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR,			

	BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG
CA 2490091	A1 20031231 CA 2003-2490091 20030619
AU 2003250847	A1 20040106 AU 2003-250847 20030619
EP 1518284	A2 20050330 EP 2003-760688 20030619
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, SK	
CN 1663064	A 20050831 CN 2003-814563 20030619
CN 100379062	C 20080402
JP 2005530676	T 20051013 JP 2004-530900 20030619
US 20060035150	A1 20060216 US 2005-518560 20050831
PRAI EP 2002-291562	A 20020621
US 2002-392978P	P 20020702
WO 2003-EP6628	W 20030619

CLASS

	PATENT NO.	CLASS	PATENT FAMILY CLASSIFICATION CODES
WO 2004001881		ICM	H01M004-58
		IPCI	H01M0004-58 [ICM, 7]
		IPCR	C01B0025-00 [I,C*]; C01B0025-45 [I,A]; C01B0025-30 [I,A]; C01B0025-37 [I,A]; C09C0003-10 [I,C*]; C09C0003-10 [I,A]; H01M0004-02 [I,C*]; H01M0004-02 [I,A]; H01M0004-36 [I,C*]; H01M0004-36 [I,A]; H01M0004-58 [I,C*]; H01M0004-58 [I,A]; H01M0004-62 [I,C*]; H01M0004-62 [I,A]; H01M0006-18 [I,C*]; H01M0006-18 [I,A]; H01M0010-36 [I,C*]; H01M0010-36 [I,A]; H01M0010-40 [I,A]
		ECLA	H01M004/58D; C01B025/37; C01B025/45; H01M004/36; H01M004/62; H01M004/62C2; H01M006/18D; H01M010/36S; T01M
CA 2490091		IPCI	H01M0004-58 [ICM, 7]; C09C0003-10 [ICS, 7]; C01B0025-30 [ICS, 7]; C01B0025-00 [ICS, 7,C*]; H01M0004-36 [ICS, 7]; H01M0004-62 [ICS, 7]
		IPCR	C01B0025-00 [I,C*]; C01B0025-45 [I,A]; C01B0025-30 [I,A]; C01B0025-37 [I,A]; C09C0003-10 [I,C*]; C09C0003-10 [I,A]; H01M0004-02 [I,C*]; H01M0004-02 [I,A]; H01M0004-36 [I,C*]; H01M0004-36 [I,A]; H01M0004-58 [I,C*]; H01M0004-58 [I,A]; H01M0004-62 [I,C*]; H01M0004-62 [I,A]; H01M0006-18 [I,C*]; H01M0006-18 [I,A]; H01M0010-36 [I,C*]; H01M0010-36 [I,A]; H01M0010-40 [I,A]
		ECLA	T01M
AU 2003250847		IPCI	H01M0004-58 [ICM, 7]
		IPCR	C01B0025-00 [I,C*]; C01B0025-45 [I,A]; C01B0025-30 [I,A]; C01B0025-37 [I,A]; C09C0003-10 [I,C*]; C09C0003-10 [I,A]; H01M0004-02 [I,C*]; H01M0004-02 [I,A]; H01M0004-36 [I,C*]; H01M0004-36 [I,A]; H01M0004-58 [I,C*]; H01M0004-58 [I,A]; H01M0004-62 [I,C*]; H01M0004-62 [I,A]; H01M0006-18 [I,C*]; H01M0006-18 [I,A]; H01M0010-36 [I,C*]; H01M0010-36 [I,A]; H01M0010-40 [I,A]
		ECLA	T01M
EP 1518284		IPCI	H01M0004-58 [ICM, 7]
		IPCR	H01M0004-58 [I,C*]; H01M0004-58 [I,A]
CN 1663064		IPCI	H01M0004-58 [I,C]; H01M0004-58 [I,A]; C01B0025-30 [I,A]; C01B0025-00 [I,C*]; C09C0003-10 [I,C]; C09C0003-10 [I,A]; H01M0004-36 [I,C]; H01M0004-36 [I,A]; H01M0004-62 [I,C]; H01M0004-62 [I,A]
		IPCR	C01B0025-00 [I,C*]; C01B0025-45 [I,A]; C01B0025-30 [I,A]; C01B0025-37 [I,A]; C09C0003-10 [I,C*]; C09C0003-10 [I,A]; H01M0004-02 [I,C*]; H01M0004-02 [I,A]; H01M0004-36 [I,C*]; H01M0004-36 [I,A]; H01M0004-58 [I,C*]; H01M0004-58 [I,A]; H01M0004-62

		[I,C*]; H01M0004-62 [I,A]; H01M0006-18 [I,C*]; H01M0006-18 [I,A]; H01M0010-36 [I,C*]; H01M0010-36 [I,A]; H01M0010-40 [I,A]
	ECLA	T01M; H01M004/58D; C01B025/37; C01B025/45; H01M004/36; H01M004/62; H01M004/62C2; H01M006/18D; H01M010/36S
JP 2005530676	IPCI	C01B0025-45 [ICM, 7]; C01B0025-00 [ICM, 7,C*]; H01M0004-02 [ICS, 7]; H01M0004-58 [ICS, 7]; H01M0010-40 [ICS, 7]; H01M0010-36 [ICS, 7,C*]
	IPCR	C01B0025-00 [I,C*]; C01B0025-45 [I,A]; C01B0025-30 [I,A]; C01B0025-37 [I,A]; C09C0003-10 [I,C*]; C09C0003-10 [I,A]; H01M0004-02 [I,C*]; H01M0004-02 [I,A]; H01M0004-36 [I,C*]; H01M0004-36 [I,A]; H01M0004-58 [I,C*]; H01M0004-58 [I,A]; H01M0004-62 [I,C*]; H01M0004-62 [I,A]; H01M0006-18 [I,C*]; H01M0006-18 [I,A]; H01M0010-36 [I,C*]; H01M0010-36 [I,A]; H01M0010-40 [I,A]
	ECLA	T01M
	FTERM	5H029/AJ02; 5H029/AJ05; 5H029/AK01; 5H029/AL12; 5H029/AM03; 5H029/AM05; 5H029/AM07; 5H029/CJ02; 5H029/CJ08; 5H029/CJ11; 5H029/CJ22; 5H029/CJ28; 5H029/DJ16; 5H029/EJ04; 5H029/HJ01; 5H029/HJ02; 5H029/HJ14; 5H050/AA02; 5H050/AA07; 5H050/BA17; 5H050/CA01; 5H050/CB12; 5H050/DA02; 5H050/DA09; 5H050/EA08; 5H050/FA17; 5H050/FA18; 5H050/GA02; 5H050/GA10; 5H050/GA11; 5H050/GA22; 5H050/GA27; 5H050/HA01; 5H050/HA02; 5H050/HA14
US 20060035150	IPCI	H01M0004-58 [I,A]; B05D0005-12 [I,A]
	IPCR	H01M0004-58 [I,A]; B05D0005-12 [I,C]; B05D0005-12 [I,A]; H01M0004-58 [I,C]
	NCL	429/221.000; 252/182.100; 427/122.000; 429/231.950
	ECLA	C01B025/45; C01B025/37; C01B025/37D; H01M004/02B; H01M004/58D; H01M004/62C2; H01M010/40L; T01M; T01M; T01M

AB The invention provides a new route for the synthesis of carbon-coated powders having the olivine or NASICON structure, which form promising classes of active products for the manufacture of rechargeable lithium batteries. Carbon-coating of the powder particles is necessary to achieve good performances because of the rather poor electronic conductivity of the structures. For the preparation of coated LiFePO<sub>4</sub>, sources of Li, Fe and phosphate are dissolved in an aqueous solution together with a polycarboxylic acid and a polyhydric alc. Upon water evaporation, polyesterification occurs while a mixed precipitate is formed containing Li, Fe and phosphate. The resin-encapsulated mixture is then heat treated at 700° in a reducing atmospheric This results in the production of a fine powder consisting of an olivine LiFePO<sub>4</sub> phase, coated with conductive carbon. When this powder is used as active material in a lithium insertion-type electrode, fast charge and discharge rates are obtained at room temperature and an excellent capacity retention is observed

ST battery carbon coated lithium contg powder prepn

IT Olivine-group minerals

RL: DEV (Device component use); USES (Uses)  
(lithium-containing; process for production of carbon-coated

lithium-containing

powders for lithium secondary battery)

IT Secondary batteries

(lithium; process for production of carbon-coated lithium-containing powders for lithium secondary battery)

IT Carboxylic acids, uses

RL: DEV (Device component use); USES (Uses)  
(polycarboxylic; process for production of carbon-coated lithium-containing powders for lithium secondary battery)

IT Polymerization  
     (polyesterification; process for production of carbon-coated lithium-containing powders for lithium secondary battery)  
 IT Alcohols, uses  
     RL: DEV (Device component use); USES (Uses)  
     (polyhydric; process for production of carbon-coated lithium-containing powders for lithium secondary battery)  
 IT Battery cathodes  
     (process for production of carbon-coated lithium-containing powders for lithium secondary battery)  
 IT 77641-62-4, Nasicon  
     RL: DEV (Device component use); USES (Uses)  
     (lithium-containing; process for production of carbon-coated lithium-containing powders for lithium secondary battery)  
 IT 10421-48-4, Ferric nitrate 13453-80-0, Lithium dihydrogen phosphate  
     RL: CPS (Chemical process); PEP (Physical, engineering or chemical process); PROC (Process)  
     (process for production of carbon-coated lithium-containing powders for lithium secondary battery)  
 IT 77-92-9, Citric acid, uses 107-21-1, Ethylene glycol,  
     uses  
     RL: DEV (Device component use); USES (Uses)  
     (process for production of carbon-coated lithium-containing powders for lithium secondary battery)  
 IT 15365-14-7P, Iron lithium phosphate felipo4  
     RL: DEV (Device component use); SPN (Synthetic preparation); PREP (Preparation); USES (Uses)  
     (process for production of carbon-coated lithium-containing powders for lithium secondary battery)  
 IT 7440-44-0, Carbon, uses  
     RL: TEM (Technical or engineered material use); USES (Uses)  
     (process for production of carbon-coated lithium-containing powders for lithium secondary battery)

L81 ANSWER 20 OF 21 CAPLUS COPYRIGHT 2008 ACS on STN  
 AN 2003:92339 CAPLUS  
 DN 138:125008  
 ED Entered STN: 06 Feb 2003  
 TI Cathode materials for secondary lithium batteries  
 IN Armand, Michel; Goodenough, John B.; Padhi, Akshaya K.; Nanjundaswamy, Kirakodu S.; Masquelier, Christian  
 PA Board of Regents, the University of Texas System, USA  
 SO U.S., 21 pp., Cont.-in-part of U.S. 5,910,382.  
     CODEN: USXXAM  
 DT Patent  
 LA English  
 IC ICM H01M004-58  
 INCL 429231100; 429218100; 429224000; 429221000  
 CC 52-2 (Electrochemical, Radiational, and Thermal Energy Technology)  
 FAN.CNT 2

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	US 6514640 US 5910382	B1 A	20030204 19990608	US 1997-998264 US 1997-840523	19971224 19970421

CA 2543784	A1	19971030	CA 1997-2543784	19970423
EP 1501137	A2	20050126	EP 2004-22447	19970423
EP 1501137	A3	20061025		
	R: DE, FR, GB, IT			
EP 1755182	A1	20070221	EP 2006-20470	19970423
	R: DE, FR, GB, IT			
EP 1755183	A1	20070221	EP 2006-21083	19970423
	R: DE, FR, GB, IT			
US 6391493	B1	20020521	US 1999-298080	19990423
US 20030082454	A1	20030501	US 2002-307346	20021202
US 20050003274	A1	20050106	US 2004-902142	20040730
US 20050244321	A1	20051103	US 2005-179617	20050713
US 20070117019	A1	20070524	US 2006-647899	20061229
US 20070166618	A1	20070719	US 2006-648316	20061229
JP 2007214147	A	20070823	JP 2007-128682	20070514
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PRAI US 1996-16060P	P	19960423		
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EP 2004-22447	A3	19970423		
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US 2002-307346	B1	20021202		
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US 2005-179617	A1	20050713		

#### CLASS

PATENT NO.	CLASS	PATENT FAMILY CLASSIFICATION CODES
US 6514640	ICM	H01M004-58
	INCL	429231100; 429218100; 429224000; 429221000
	IPCI	H01M0004-58 [ICM, 7]
	IPCR	C01B0025-00 [I,C*]; C01B0025-45 [I,A]; H01M0004-02 [N,C*]; H01M0004-02 [N,A]; H01M0004-58 [I,C*]; H01M0004-58 [I,A]; H01M0010-36 [N,C*]; H01M0010-40 [N,A]
	NCL	429/231.100; 429/218.100; 429/221.000; 429/224.000
	ECLA	C01B025/45; H01M004/58D; T01M; T01M; T01M
US 5910382	IPCI	H01M0004-58 [ICM, 6]
	IPCR	C01B0025-00 [I,C*]; C01B0025-45 [I,A]; H01M0004-02 [N,C*]; H01M0004-02 [N,A]; H01M0004-58 [I,C*]; H01M0004-58 [I,A]; H01M0010-36 [N,C*]; H01M0010-40 [N,A]
	NCL	429/218.100; 429/221.000; 429/224.000
	ECLA	C01B025/45; H01M004/58D; T01M; T01M; T01M
CA 2543784	IPCI	H01M0004-40 [I,A]; H01M0010-00 [I,A]
	IPCR	H01M0004-40 [I,A]; H01M0004-40 [I,C]; H01M0010-00 [I,C]; H01M0010-00 [I,A]
EP 1501137	IPCI	H01M0004-58 [I,A]; C01B0025-26 [I,A]; C01B0025-45 [I,A]; C01B0025-00 [I,C*]
	ECLA	H01M004/58D; C01B025/45; T01M; T01M
EP 1755182	IPCI	H01M0004-58 [I,A]; C01B0025-26 [I,A]; C01B0025-45 [I,A]; C01B0025-00 [I,C*]
	ECLA	C01B025/45; H01M004/58D; T01M; T01M; T01M; T01M
EP 1755183	IPCI	H01M0004-58 [I,A]; C01B0025-26 [I,A]; C01B0025-45 [I,A]; C01B0025-00 [I,C*]
	ECLA	C01B025/45; H01M004/58D; T01M; T01M; T01M; T01M
US 6391493	IPCI	H01M0004-58 [ICM, 7]
	IPCR	C01B0025-00 [I,C*]; C01B0025-45 [I,A]; H01M0004-02 [N,C*]; H01M0004-02 [N,A]; H01M0004-58 [I,C*];

		H01M0004-58 [I,A]; H01M0010-36 [N,C*]; H01M0010-40 [N,A]
	NCL	429/218.100; 429/231.100
	ECLA	C01B025/45; H01M004/58D; T01M; T01M; T01M; T01M
US 20030082454	IPCI	H01M0004-58 [ICM,7]
	IPCR	C01B0025-00 [I,C*]; C01B0025-45 [I,A]; H01M0004-02 [N,C*]; H01M0004-02 [N,A]; H01M0004-58 [I,C*]; H01M0004-58 [I,A]; H01M0010-36 [N,C*]; H01M0010-40 [N,A]
	NCL	429/231.950; 429/221.000; 429/223.000; 429/224.000; 429/231.500
US 20050003274	ECLA	C01B025/45; H01M004/58D; T01M; T01M; T01M; T01M
	IPCI	H01M0004-58 [ICM,7]; H01M0004-62 [ICS,7]; C01B0033-32 [ICS,7]; C01B0033-00 [ICS,7,C*]; C01B0025-26 [ICS,7]; C01B0025-00 [ICS,7,C*]; C01B0017-98 [ICS,7]; C01B0017-00 [ICS,7,C*]; C01G0031-02 [ICS,7]; C01G0031-00 [ICS,7,C*]
	IPCR	C01B0025-00 [I,C*]; C01B0025-45 [I,A]; H01M0004-02 [N,C*]; H01M0004-02 [N,A]; H01M0004-58 [I,C*]; H01M0004-58 [I,A]; H01M0010-36 [N,C*]; H01M0010-40 [N,A]
	NCL	429/231.950; 423/306.000; 423/332.000; 423/518.000; 423/594.800; 429/217.000; 429/220.000; 429/221.000; 429/223.000; 429/224.000; 429/229.000; 429/231.500
US 20050244321	ECLA	C01B025/45; H01M004/58D; T01M; T01M; T01M; T01M
	IPCI	H01M0004-58 [ICM,7]; C01B0025-45 [ICS,7]; C01B0025-00 [ICS,7,C*]
	IPCR	C01B0025-00 [I,C*]; C01B0025-45 [I,A]; H01M0004-58 [I,C*]; H01M0004-58 [I,A]; H01M0010-36 [N,C*]; H01M0010-40 [N,A]
	NCL	423/306.000; 252/182.100; 429/221.000; 429/223.000; 429/224.000; 429/231.500; 429/231.900; 429/231.950
US 20070117019	ECLA	H01M004/58D; C01B025/45; T01M
	IPCI	H01M0004-58 [I,A]; C01B0025-45 [I,A]; C01B0025-00 [I,C*]
	IPCR	H01M0004-58 [I,C]; H01M0004-58 [I,A]; C01B0025-00 [I,C]; C01B0025-45 [I,A]
	NCL	429/231.950; 423/306.000; 429/221.000; 429/223.000; 429/224.000; 429/231.500
US 20070166618	ECLA	C01B025/45
	IPCI	H01M0004-58 [I,A]; C01B0025-45 [I,A]; C01B0025-00 [I,C*]
	IPCR	H01M0004-58 [I,C]; H01M0004-58 [I,A]; C01B0025-00 [I,C]; C01B0025-45 [I,A]
	NCL	429/231.950; 423/306.000; 429/221.000; 429/223.000; 429/224.000
	ECLA	C01B025/45
JP 2007214147	IPCI	H01M0004-58 [I,A]
	IPCR	H01M0004-58 [I,C]; H01M0004-58 [I,A]; C01B0025-00 [I,C*]; C01B0025-45 [I,A]; H01M0004-02 [I,C*]; H01M0004-02 [I,A]; H01M0010-36 [N,C*]; H01M0010-40 [N,A]
	ECLA	C01B025/45; H01M004/58D; T01M; T01M; T01M; T01M
	FTERM	5H050/AA02; 5H050/BA17; 5H050/CA01; 5H050/CB12; 5H050/GA02; 5H050/GA10; 5H050/GA12; 5H050/GA27; 5H050/GA28; 5H050/HA02; 5H050/HA14; 5H050/HA17; 5H050/HA18; 5H050/HA19
JP 2007294463	IPCI	H01M0004-58 [I,A]
	IPCR	H01M0004-58 [I,C]; H01M0004-58 [I,A]; C01B0025-00 [I,C*]; C01B0025-45 [I,A]; H01M0004-02 [I,C*]; H01M0004-02 [I,A]; H01M0010-36 [N,C*]; H01M0010-40 [N,A]

ECLA C01B025/45; H01M004/58D; T01M; T01M; T01M; T01M  
 FTERM 5H050/AA01; 5H050/AA02; 5H050/AA17; 5H050/BA16;  
 5H050/BA17; 5H050/CA01; 5H050/CB12; 5H050/FA17;  
 5H050/HA02; 5H050/HA05; 5H050/HA18  
 US 20070281215 IPCI H01M0004-58 [I,A]; C01B0025-45 [I,A]; C01B0025-00  
 [I,C\*]  
 IPCR H01M0004-58 [I,C]; H01M0004-58 [I,A]; C01B0025-00  
 [I,C]; C01B0025-45 [I,A]  
 NCL 429/231.950; 423/306.000; 429/221.000; 429/223.000;  
 429/224.000; 429/231.500  
 ECLA C01B025/45

**AB** The invention relates to materials for use as electrodes in an alkali-ion secondary battery, particularly a lithium-ion battery. The invention provides transition-metal compds. having the ordered-olivine, a modified olivine, or the rhombohedral NASICON structure and the polyanion (PO<sub>4</sub>)<sub>3</sub><sup>-</sup> as at least one constituent for use as electrode material for alkali-ion rechargeable batteries.

**ST** cathode lithium secondary battery  
**IT** Transition metal nitrides  
 RL: DEV (Device component use); USES (Uses)  
 (Li-containing; cathode materials for secondary lithium batteries)

**IT** EPDM rubber  
 Fluoropolymers, uses  
 Polyesters, uses  
 Polyethers, uses  
 RL: MOA (Modifier or additive use); USES (Uses)  
 (binder; cathode materials for secondary lithium batteries)

**IT** Battery anodes  
 Battery cathodes  
 (cathode materials for secondary lithium batteries)

**IT** Chalcogenides  
 RL: DEV (Device component use); USES (Uses)  
 (lamellar; cathode materials for secondary lithium batteries)

**IT** Secondary batteries  
 (lithium; cathode materials for secondary lithium batteries)

**IT** Lithium alloy, base  
 RL: DEV (Device component use); USES (Uses)  
 (cathode materials for secondary lithium batteries)

**IT** 116-14-3D, Tetrafluoroethylene, copolymer 9002-84-0, Ptfe 9011-14-7,  
 Pmma 24937-79-9, Pvdf 25014-41-9, Polyacrylonitrile  
 RL: MOA (Modifier or additive use); USES (Uses)  
 (binder; cathode materials for secondary lithium batteries)

**IT** 69104-84-3, Sodium vanadiumphosphate Na<sub>3</sub>V<sub>2</sub>(PO<sub>4</sub>)<sub>3</sub>  
 RL: CPS (Chemical process); PEP (Physical, engineering or chemical process); PROC (Process)  
 (cathode materials for secondary lithium batteries)

**IT** 96-48-0,  $\gamma$ -Butyrolactone 96-49-1, Ethylene carbonate 105-58-8,  
 Diethyl carbonate 107-21-1D, Ethylene glycol, dialkyl ether  
 108-32-7, Propylene carbonate 111-46-6D, DiEthylene glycol,  
 dialkyl ether 112-27-6D, TriEthylene glycol, dialkyl ether 112-60-7D,  
 TetraEthylene glycol, dialkyl ether 616-38-6, Dimethyl carbonate  
 623-53-0, Methyl ethyl carbonate 7439-93-2, Lithium, uses 7803-58-9D,  
 Sulfamide, tetraalkyl derivative 36058-25-0, Iron lithium phosphate  
 Fe<sub>2</sub>Li<sub>3</sub>(PO<sub>4</sub>)<sub>3</sub> 39302-37-9, Lithium titanium oxide 39448-96-9,  
 Graphite-lithium 77641-62-4, Nasicon 223505-09-7, Iron lithium  
 titanium phosphate 277742-93-5, Vanadium oxide VO<sub>2.1-2.5</sub>  
 RL: DEV (Device component use); USES (Uses)  
 (cathode materials for secondary lithium batteries)

**IT** 13824-63-0P, Cobalt lithium phosphate colipo<sub>4</sub> 13826-59-0P, Lithium  
 manganese phosphate limnpo<sub>4</sub> 13977-83-8P, Lithium nickel phosphate  
 linipo<sub>4</sub> 15365-14-7P, Iron lithium phosphate felipo<sub>4</sub>

37144-98-2P, Niobium titanium phosphate NbTi(PO<sub>4</sub>)<sub>3</sub> 161774-31-8P, Iron lithium niobium phosphate FeLiNb(PO<sub>4</sub>)<sub>3</sub> 184241-62-1P 205380-60-5P, Iron lithium phosphate sulfate Fe<sub>2</sub>Li(PO<sub>4</sub>)(SO<sub>4</sub>)<sub>2</sub> 488829-05-6P, Iron lithium manganese phosphate (Fe0.5-LiMn0-0.5(PO<sub>4</sub>)) 488829-06-7P, Iron lithium titanium phosphate silicate (Fe0.8Li1.1Ti(PO<sub>4</sub>)<sub>0.8</sub>(SiO<sub>4</sub>)<sub>0.2</sub>) 951777-58-5P, Lithium sodium vanadium phosphate Li<sub>2</sub>NaV<sub>2</sub>(PO<sub>4</sub>)<sub>3</sub>  
 RL: DEV (Device component use); SPN (Synthetic preparation); PREP (Preparation); USES (Uses)  
 (cathode materials for secondary lithium batteries)  
 IT 7440-44-0, Carbon, uses  
 RL: MOA (Modifier or additive use); USES (Uses)  
 (cathode materials for secondary lithium batteries)  
 RE.CNT 9 THERE ARE 9 CITED REFERENCES AVAILABLE FOR THIS RECORD  
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 AN 2001:796403 CAPLUS  
 DN 135:346864  
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 TI Cathode for nonaqueous electrolyte lithium ion battery  
 IN Yamada, Atsuo; Yamahira, Takayuki  
 PA Sony Corporation, Japan  
 SO Eur. Pat. Appl., 26 pp.  
 CODEN: EPXXDW  
 DT Patent  
 LA English  
 IC ICM H01M004-58  
 ICS C01G049-00; C01B025-30; C01B025-45; H01M004-38  
 CC 52-2 (Electrochemical, Radiational, and Thermal Energy Technology)  
 FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	EP 1150368	A2	20011031	EP 2001-109919	20010424
	EP 1150368	A3	20051026		
	R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO				
	JP 2001307730	A	20011102	JP 2000-128998	20000425
	JP 3959929	B2	20070815		
	MX 2001PA04029	A	20030820	MX 2001-PA4029	20010423
	TW 533617	B	20030521	TW 2001-90109790	20010424
	CA 2344981	A1	20011025	CA 2001-2344981	20010425
	CN 1320976	A	20011107	CN 2001-117211	20010425
	US 20020004169	A1	20020110	US 2001-842485	20010425
	US 6746799	B2	20040608		
PRAI	JP 2000-128998	A	20000425		

#### CLASS

PATENT NO.	CLASS	PATENT FAMILY CLASSIFICATION CODES
EP 1150368	ICM	H01M004-58
	ICS	C01G049-00; C01B025-30; C01B025-45; H01M004-38
	IPCI	H01M0004-58 [ICM,6]; C01G0049-00 [ICS,6]; C01B0025-30 [ICS,6]; C01B0025-45 [ICS,6]; C01B0025-00 [ICS,6,C*]; H01M0004-38 [ICS,6]

	IPCR	H01M0010-36 [I,C*]; H01M0010-40 [I,A]; C01B0025-00 [I,C*]; C01B0025-45 [I,A]; H01M0004-02 [I,C*]; H01M0004-02 [I,A]; H01M0004-36 [I,C*]; H01M0004-36 [I,A]; H01M0004-38 [I,C*]; H01M0004-38 [I,A]; H01M0004-40 [N,C*]; H01M0004-40 [N,A]; H01M0004-48 [N,C*]; H01M0004-48 [N,A]; H01M0004-52 [I,C*]; H01M0004-52 [I,A]; H01M0004-58 [I,C*]; H01M0004-58 [I,A]; H01M0010-36 [N,A]
	ECLA	H01M004/525; C01B025/45; H01M004/131; H01M004/38; H01M004/58D; T01M; T01M; T01M
JP 2001307730	IPCI	H01M0004-58 [I,A]; H01M0004-02 [I,A]; H01M0010-40 [I,A]; H01M0010-36 [I,C*]
	IPCR	H01M0010-36 [I,C*]; H01M0010-40 [I,A]; C01B0025-00 [I,C*]; C01B0025-45 [I,A]; H01M0004-02 [I,C*]; H01M0004-02 [I,A]; H01M0004-36 [I,C*]; H01M0004-36 [I,A]; H01M0004-38 [I,C*]; H01M0004-38 [I,A]; H01M0004-40 [N,C*]; H01M0004-40 [N,A]; H01M0004-48 [N,C*]; H01M0004-48 [N,A]; H01M0004-52 [I,C*]; H01M0004-52 [I,A]; H01M0004-58 [I,C*]; H01M0004-58 [I,A]; H01M0010-36 [N,A]
MX 2001PA04029	IPCI	H01M0004-50 [ICM,7]; H01M0004-58 [ICS,7]
TW 533617	IPCI	H01M0004-64 [ICM,7]
	IPCR	H01M0010-36 [I,C*]; H01M0010-40 [I,A]; C01B0025-00 [I,C*]; C01B0025-45 [I,A]; H01M0004-02 [I,C*]; H01M0004-02 [I,A]; H01M0004-36 [I,C*]; H01M0004-36 [I,A]; H01M0004-38 [I,C*]; H01M0004-38 [I,A]; H01M0004-40 [N,C*]; H01M0004-40 [N,A]; H01M0004-48 [N,C*]; H01M0004-48 [N,A]; H01M0004-52 [I,C*]; H01M0004-52 [I,A]; H01M0004-58 [I,C*]; H01M0004-58 [I,A]; H01M0010-36 [N,A]
CA 2344981	IPCI	H01M0004-36 [ICM,7]; H01M0004-24 [ICS,7]; H01M0010-24 [ICS,7]
	IPCR	H01M0010-36 [I,C*]; H01M0010-40 [I,A]; C01B0025-00 [I,C*]; C01B0025-45 [I,A]; H01M0004-02 [I,C*]; H01M0004-02 [I,A]; H01M0004-36 [I,C*]; H01M0004-36 [I,A]; H01M0004-38 [I,C*]; H01M0004-38 [I,A]; H01M0004-40 [N,C*]; H01M0004-40 [N,A]; H01M0004-48 [N,C*]; H01M0004-48 [N,A]; H01M0004-52 [I,C*]; H01M0004-52 [I,A]; H01M0004-58 [I,C*]; H01M0004-58 [I,A]; H01M0010-36 [N,A]
CN 1320976	IPCI	H01M0004-48 [ICM]; H01M0010-36 [ICS]
	IPCR	H01M0010-36 [I,C*]; H01M0010-40 [I,A]; C01B0025-00 [I,C*]; C01B0025-45 [I,A]; H01M0004-02 [I,C*]; H01M0004-02 [I,A]; H01M0004-36 [I,C*]; H01M0004-36 [I,A]; H01M0004-38 [I,C*]; H01M0004-38 [I,A]; H01M0004-40 [N,C*]; H01M0004-40 [N,A]; H01M0004-48 [N,C*]; H01M0004-48 [N,A]; H01M0004-52 [I,C*]; H01M0004-52 [I,A]; H01M0004-58 [I,C*]; H01M0004-58 [I,A]; H01M0010-36 [N,A]
US 20020004169	IPCI	H01M0004-58 [ICM,7]
	IPCR	H01M0010-36 [I,C*]; H01M0010-40 [I,A]; C01B0025-00 [I,C*]; C01B0025-45 [I,A]; H01M0004-02 [I,C*]; H01M0004-02 [I,A]; H01M0004-36 [I,C*]; H01M0004-36 [I,A]; H01M0004-40 [N,C*]; H01M0004-40 [N,A]; H01M0004-48 [N,C*]; H01M0004-48 [N,A]; H01M0004-58 [I,C*]; H01M0004-58 [I,A]; H01M0004-38 [I,C*]; H01M0004-38 [I,A]; H01M0004-52 [I,C*]; H01M0004-52 [I,A]; H01M0010-36 [N,A]
	NCL	429/221.000; 429/224.000; 429/231.100; 429/231.800; 429/218.100
	ECLA	H01M004/525; C01B025/45; H01M004/131; H01M004/38; H01M004/58D; T01M; T01M; T01M

AB The lithium ion cell is improved appreciably in operational stability under special conditions, such as high temps., and exhibits superior characteristics against over-discharging, while guaranteeing compatibility to the operating voltage of a conventional lithium ion cell and an energy d. equivalent to that of the conventional lithium ion cell. To this end, the lithium ion cell includes a pos. electrode, a neg. electrode and a nonaq. electrolyte, and uses, as a pos. electrode active material, a composite material of a first lithium compound represented by the general formula  $LixMyPO_4$ , where  $0 < x < 2$ ,  $0.8 < y < 1.2$  and M contains Fe, and a second lithium compound having a potential holder than the potential of the first lithium compound

ST lithium nonaq electrolyte cathode

IT Charcoal

RL: DEV (Device component use); USES (Uses)  
(activated; cathode for nonaq. electrolyte lithium ion battery)

IT Battery cathodes  
(cathode for nonaq. electrolyte lithium ion battery)

IT Carbon fibers, uses  
Carbonaceous materials (technological products)

Coke  
Petroleum coke

RL: DEV (Device component use); USES (Uses)  
(cathode for nonaq. electrolyte lithium ion battery)

IT Carbon black, uses  
RL: MOA (Modifier or additive use); USES (Uses)  
(cathode for nonaq. electrolyte lithium ion battery)

IT Fluoropolymers, uses  
RL: TEM (Technical or engineered material use); USES (Uses)  
(cathode for nonaq. electrolyte lithium ion battery)

IT Organic compounds, uses  
RL: DEV (Device component use); USES (Uses)  
(high mol., sintered; cathode for nonaq. electrolyte lithium ion battery)

IT Secondary batteries  
(lithium; cathode for nonaq. electrolyte lithium ion battery)

IT Coke  
RL: DEV (Device component use); USES (Uses)  
(needle; cathode for nonaq. electrolyte lithium ion battery)

IT Coke  
RL: DEV (Device component use); USES (Uses)  
(pitch; cathode for nonaq. electrolyte lithium ion battery)

IT Furan resins  
Phenolic resins, uses

RL: DEV (Device component use); USES (Uses)  
(sintered and carbonized; cathode for nonaq. electrolyte lithium ion battery)

IT 50-21-5D, Lactic acid, ester 60-29-7, Diethyl ether, uses  
64-19-7D, Acetic acid, ester, uses 75-05-8, Acetonitrile, uses  
79-09-4D, Propionic acid, ester 96-47-9, 2-Methyltetrahydrofuran  
96-48-0 96-49-1, Ethylene carbonate 100-66-3, Anisole, uses  
105-58-8, Diethyl carbonate 107-12-0, Propionitrile 108-32-7,  
Propylene carbonate 109-99-9, Thf, uses 110-71-4, 1,2-Dimethoxyethane  
126-33-0, Sulfolane 409-21-2, Silicon carbide sic, uses 554-12-1,  
Methyl propionate 616-38-6, Dimethyl carbonate 623-42-7, Methyl  
butyrate 623-96-1, Dipropyl carbonate 629-14-1, 1,2-Diethoxyethane  
646-06-0, 1,3-Dioxolane 872-36-6, Vinylene carbonate 1072-47-5,  
4-Methyl-1,3-dioxolane 1313-08-2 2550-62-1, Lithium methanesulfonate  
4437-85-8, Butylene carbonate 7439-93-2, Lithium, uses 7440-50-8,  
Copper, uses 7447-41-8, Lithium chloride, uses 7550-35-8, Lithium  
bromide 7782-42-5, Graphite, uses 7791-03-9, Lithium perchlorate  
9003-07-0, Polypropylene 12007-81-7, Silicon tetraboride 12008-29-6,

Silicon hexaboride 12013-56-8, Calcium disilicide 12017-12-8, Cobalt disilicide 12018-09-6, Chromium disilicide 12022-99-0, Iron disilicide 12032-86-9, Manganese disilicide 12033-76-0, Silicon nitride oxide Si<sub>2</sub>N<sub>2</sub>O 12033-89-5, Silicon nitride, uses 12034-80-9, Niobium disilicide 12039-79-1, Tantalum disilicide 12039-83-7, Titanium silicide TiSi<sub>2</sub> 12039-87-1, Vanadium disilicide 12039-88-2, Tungsten disilicide 12059-14-2, Nickel silicide (Ni<sub>2</sub>Si) 12136-78-6, Molybdenum disilicide 12159-07-8, Copper silicide Cu<sub>5</sub>Si 12190-79-3, Cobalt lithium oxide Colio<sub>2</sub> 12201-89-7, Nickel disilicide 14283-07-9, Lithium tetrafluoroborate 14485-20-2, Lithium tetraphenylborate 15365-14-7, Iron lithium phosphate FeLiPO<sub>4</sub> 19414-36-9, Iron lithium manganese phosphate ((Fe,Mn)Li(PO<sub>4</sub>)) 21324-40-3, Lithium hexafluorophosphate 22831-39-6, Magnesium silicide (Mg<sub>2</sub>Si) 29935-35-1, Lithium hexafluoroarsenate 33454-82-9, Lithium trifluoromethanesulfonate 35678-71-8, Methylsulfolane 90076-65-6 113066-89-0, Cobalt lithium nickel oxide Co<sub>0.2</sub>LiNiO<sub>0.8</sub>O<sub>2</sub> 113671-38-8, Silicon oxide SiO<sub>0.2</sub> 160479-36-7, Lithium tin oxide 178958-56-0, Lithium silicon oxide 300858-61-1 339333-78-7, Zinc silicide ZnSi<sub>2</sub> 371148-86-6, Tin oxide (SnO<sub>0.2</sub>) 371148-87-7, Lithium magnesium manganese oxide (LiMg<sub>0.2</sub>Mn<sub>0.8</sub>O<sub>2</sub>) RL: DEV (Device component use); USES (Uses)  
(cathode for nonaq. electrolyte lithium ion battery)

IT 24937-79-9, Pvdf  
RL: TEM (Technical or engineered material use); USES (Uses)  
(cathode for nonaq. electrolyte lithium ion battery)

IT 7440-44-0, Carbon, uses  
RL: DEV (Device component use); USES (Uses)  
(pyrocarbon; cathode for nonaq. electrolyte lithium ion battery)

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COST IN U.S. DOLLARS          SINCE FILE      TOTAL
                                ENTRY        SESSION
FULL ESTIMATED COST          85.51         203.57

DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS) SINCE FILE      TOTAL
                                              ENTRY        SESSION
CA SUBSCRIBER PRICE           -16.80         -17.60
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